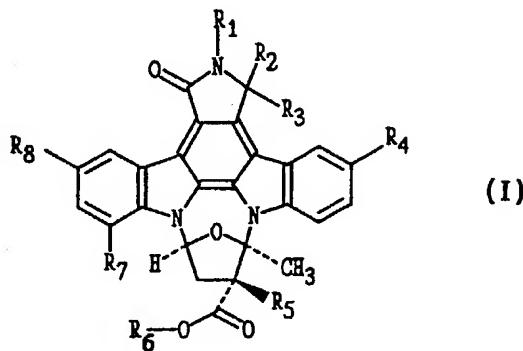




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(22) International Filing Date: 30 July 1993 (30.07.93)		(75) Inventors/Applicants (for US only) : ABRAHAM, Irene [US/US]; 1805 Waite, Kalamazoo, MI 49008 (US). ARISTOFF, Paul, A. [US/US]; 1650 Brookmoor Lane, Portage, MI 49002 (US). SKULNICK, Harvey, I. [US/US]; 1745 Old Deer Run, Kalamazoo, MI 49009 (US).	
(30) Priority data: 07/929,199 12 August 1992 (12.08.92) US		(74) Agent: WOOTTON, Thomas, A.; The Upjohn Company, 301 Henrietta Street, Kalamazoo, MI 49001 (US).	
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(71) Applicant (for all designated States except US): THE UPJOHN COMPANY [US/US]; 301 Henrietta Street, Kalamazoo, MI 49001 (US).		Published <i>Without international search report and to be republished upon receipt of that report.</i>	

(54) Title: PROTEIN KINASE INHIBITORS AND RELATED COMPOUNDS COMBINED WITH TAXOL



(57) Abstract

This invention describes both known and novel compounds, some of which are protein kinase inhibitors, that may be combined with taxol type compounds. The combination of disclosed compounds plus taxol type compounds exhibits powerful synergistic effects and the combinations are useful in the treatment of cancer. The novel compounds and their synthesis are described. A compound of formula (I), above, is described wherein R₁-R₈ represent various substituents.

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PROTEIN KINASE INHIBITORS AND RELATED
COMPOUNDS COMBINED WITH TAXOL
FIELD OF THE INVENTION

This invention describes the use of compounds that are used in combination with taxol
5 to control cancerous growths and tumors. Protein kinase inhibitors and related compounds are
combined with taxol and taxol related compounds and the combination of compounds exhibits
powerful potentiating effects when used to treat cancer. Many of the compounds are protein
kinase inhibitors, other compounds achieve similar effects but are not necessarily protein kinase
inhibitors.

10 **BACKGROUND OF THE INVENTION**

Taxol was first isolated from the bark of the western yew, *Taxus brevifolia*, and
identified as an antitumor agent in 1971 by Wani, et al. Recently, phase II clinical trials with
taxol have shown it to be one of the most exciting chemotherapeutics available. Taxol has
proven effective in drug-refractory ovarian cancer (McGuire, et al., 1989), and has shown a 56%
15 objective response rate in metastatic breast cancer (Holmes, et al., 1991). In addition, there is
reason for hope that taxol may be effective in many other types of cancers.

The development of taxol, however, has faced many obstacles. Taxol's poor solubility
required that it be administered in the vehicle Cremophor EL (polyethylated castor oil), which
led to a high incidence of hypersensitivity reactions. It is not clear whether these reactions are
20 caused by the vehicle or the drug, but it was found that using longer drug infusions (Weiss, et
al., 1990) and anti-allergic regimens (Rowinsky, et al., 1990) reduced the incidence of such
reactions. In addition, there are inherent problems in producing sufficient quantities of taxol.
Extraction from the bark of the extremely slow growing western yew using present methods
cannot meet the demand for taxol. Cultivation of the western yew may take years to establish,
25 synthesis of the complex taxol molecule will be difficult and/or very expensive. Alternative
sources of taxol or a taxol substitute or a taxol additive would therefore be highly desirable.

Taxol has been shown previously to be toxic to tumor and leukemia cells inoculated in
mice, including L-1210, P388 and P-1534 leukemia cells and Walker WM-256 carcinosarcoma,
sarcoma 180 and Lewis lung tumor cells (Wani et al., 1971). It has also been shown to be toxic
30 to cultured human HeLa cells (Schiff et al., 1979) and CHO (Chinese hamster ovary) cells
(Cabral et al., 1981). This evidence of toxicity to rodent and human tumor cells *in vitro* and to
tumor bearing mice *in vivo* predicted that taxol would be an active chemotherapeutic agent and
led to clinical trials in human cancer patients. These clinical trials showed efficacy of taxol in
treating ovarian cancer (McGuire et al., 1989). Taxotere is a taxol type compound that has also
35 been shown to have powerful antitumor activity. Bissery et. al, Cancer Research 51, 4845-4852,
Sept. 15, 1991.

-2-

Since taxol is now known to be an effective chemotherapeutic agent, a co-treatment that increases the toxicity of taxol on cancer or transformed cells, such as CHO cells, would be likely to increase the chemotherapeutic effect of taxol in cancer patients or to allow smaller doses of taxol to be administered. Quantities of taxol available are extremely limited.

5 Compounds that increase the efficacy of taxol, thereby allowing smaller amounts to be used with equal effectiveness, will enable more patients to be treated with taxol. This should also reduce the hypersensitivity and non-therapeutic toxic reactions seen clinically with taxol, as both less taxol and the less of the vehicle needed to deliver taxol will need to be administered.

Our finding is that when the compounds of this invention are combined with taxol or
10 taxol related compounds the mixture of compounds has a potentiating effect that, surprisingly produces tumor cell toxicity at lower doses than taxol alone. These findings, and other studies, suggest that the compounds will be effective in synergizing with taxol in killing tumor cells in human cancer patients. The findings also suggest results may be seen with taxol related compounds such as taxotere and related taxol analogues.

15 **INFORMATION DISCLOSURE**

Many of the compounds of this invention are related to the physiologically active substance K-252. The following patents disclose some of these compounds. U.S. 4,877,776 issued October 31, 1989. U.S. 4,923,986 issued May 8, 1990. W.O 8807-045-A published September 22, 1988.

20 The following Japanese patent applications also disclose related compounds: J63 295-588-A, J63 295-589-A, J62 155-284-A, and J62 155-285-A disclose stausporine related compounds.

SUMMARY OF THE INVENTION

This invention is in two parts. Known compounds are listed in part I, they are claimed
25 for the method of using the compounds as described herein. The known compounds are also claimed as compounds combined in a composition with taxol type compounds. The new compounds are in part II. The new compounds are claimed as compounds, for their method of use and in a composition.

I. The known compounds.

30 A. Indolocarbazole Type Compounds

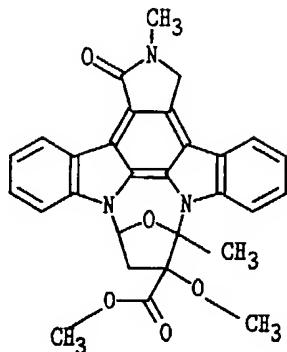
1) The "First Known Derivatives of K-252." The "First Known Derivatives of K-252" are all of the compounds disclosed in U.S. patent 4,877,776. U.S. patent 4,877,776 incorporated herein by reference.

2) The "Second Known Derivatives of K-252." The "Second Known Derivatives of K-252" are all of the compounds disclosed in U.S. patent 4,923,986. U.S. patent 4,923,986 incorporated herein by reference.

-3-

3) The specific compounds below are more preferred,
3(a) KT5823

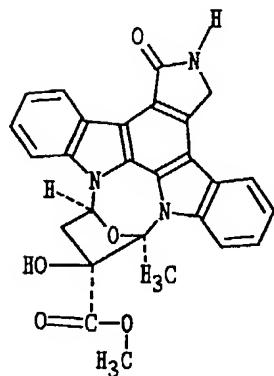
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3(b) K-252A

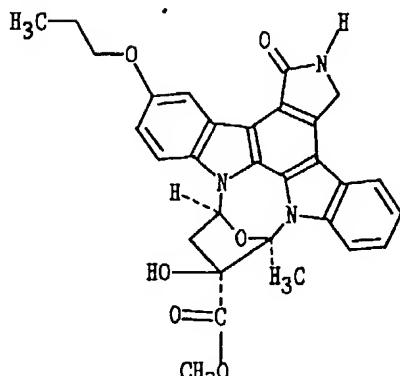
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3(c) KT5926

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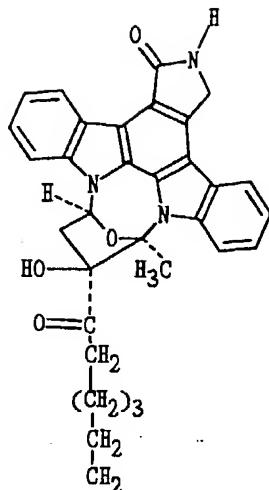
-4-

3(d) KT5720

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3(e) Staurosporine

B. Non-Indolecarbazole Type Compounds

- 1) Adriamycin
- 2) Amiloride
- 3) Calphostin
- 4) Chlorpromazine
- 5) The compound known as "HA-1004"
- 6) Indomethacin
- 7) Okadaic acid
- 8) Phenazocine
- 9) Polymyxin B
- 10) 2-aminopurine
- 11) 6-dimethyl-aminopurine
- 12) Sphingosine
- 13) Tamoxifen
- 14) Compounds related to tamoxifen such as triphenylethylene antiestrogens
- 15) Trifluoperazine
- 16) Verapamil
- 17) 3-isobutyl-1-methyl-xanthine
- 18) 8-Cl-cAMP

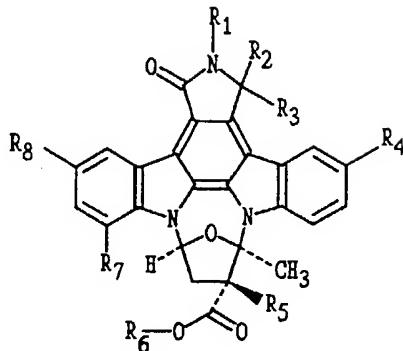
-5-

II. The new compounds.

A compound of FORMULA I, below,

5

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FORMULA I

wherein,

R₁ is -H, -(C₁-C₄ alkyl), -C(O)-(C₁-C₄ alkyl), -NH₂,
 -C(O)-NH₂, -CH₂CH₂-N(R₁₋₁)₂,

15 wherein R₁₋₁ is -H or -(C₁-C₄) alkyl,

R₂ is -H, or R₂ and R₃ taken together are (O),

R₃ is -H, -OH or R₂ and R₃ taken together are (O),

R₄ is -H, -OH, -NH₂, or -O-(C₁-C₄ alkyl),

R₅ is -OH, -O-(C₁-C₄ alkyl), or -O-C(O)-(C₁-C₄ alkyl),

20 R₆ is -(C₆-C₁₂ alkyl), -(C₃-C₁₀ cycloalkyl), -(CH₂)_nCH₂N(R₆₋₁)₂,

wherein R₆₋₁ is -H, or -(C₁-C₄) alkyl,

R₇ is -H, or -NH₂,

R₈ is -Cl, -Br, -H, -CH₃, -CH₂OH, -OH; -O-(C₁-C₄ alkyl),

-N(R₈₋₁)₂, or -NHC(O)-NH(R₈₋₁),

25 wherein R₈₋₁ is -H or -(C₁-C₄ alkyl)

wherein n is 0-5

with the proviso that:

a) when R₂ or R₃ is -OH then R₁ is H,

b) when R₁, R₂, R₃, R₄, and R₇ all equal H and R₅ is OH

30 then R₆ does NOT equal -(CH₂)₅CH₃,

c) when R₁, R₄, and R₇ all equal H, and R₂ combined with R₃ is (O),

and R₅ is OH, then R₆ does NOT equal -(CH₂)₅CH₃,

d) when R₄ is -OH, -NH₂, or -O(C₁-C₄ alkyl), then R₄ and R₈ are the same.

35 A pharmaceutical composition consisting of a pharmaceutically acceptable carrier and an effective amount of FORMULA I. A pharmaceutical composition consisting of a

pharmaceutically acceptable carrier and an effective amount of the compound of FORMULA I in conjunction with an appropriate dose of taxol or taxol related compounds. A method of controlling cancerous growths in mammals which comprises administering a therapeutic or prophylactic dosage of any of the three following groups of compounds in conjunction with an appropriate dose of taxol or taxol related compounds. 1) a compound of FORMULA I, 2) any one of the compounds described in the specification as "Indolecarbazole Type Compounds.", 3) any one of the compounds described in the specification as "Non-Indolecarbazole Type Compounds."

BRIEF DESCRIPTION OF THE FIGURES

10 Figure 1. Isobogram showing potentiating effect of the combination of Taxol plus KT5823. The isobogram shows the effectiveness of a combination of 2 drugs for the killing of wild type, 10001a CHO, cells. The data line is the solid line with open circle or triangle data points. The data line shows the combination of doses which gives an LD₅₀ for the cells. The diagonal dashed line shows the predicted concentrations of drugs if their combination only had an additive effect. If any data points were above the dashed line that would indicate the combination of compounds had antagonistic effects. Data points below the line indicate the compounds have potentiating or synergistic effects.

Figure 2. Isobogram showing potentiating effect of the combination of taxol plus KT5926.

20 Figure 3. Isobogram showing potentiating effect of the combination of taxol plus KT5720.

Figure 4. Isobogram showing NO potentiating effect from the combination of taxol plus H-9. This isobogram shows the predicted effect of a "control" substance that does NOT act in a potentiating or synergistic manner.

25 Figure 5. Isobogram showing potentiating effect of the combination of taxol plus K252a.

Figure 6. Isobogram showing potentiating effect of the combination of taxol plus tamoxifen.

30 Figure 7. Isobogram showing potentiating effect of the combination of taxol plus 2-aminopurine.

Figure 8. Isobogram showing potentiating effect of the combination of taxol plus 6-dimethylaminopurine.

Figure 9. Isobogram showing potentiating effect of the combination of taxol plus chlorpromazine.

35 Figure 10. Isobogram showing potentiating effect of the combination of taxol plus 3-isobutyl-1-methyl-xanthine.

Figure 11. Isobologram showing potentiating effect of the combination of taxol plus 8-Cl-cAMP.

Figure 12. Isobologram showing potentiating effect of the combination of taxol plus Example A-1.

5 Figure 13. Isobologram showing potentiating effect of the combination of taxol plus Example B-1.

Figure 14. Isobologram showing potentiating effect of the combination of taxol plus Example B-2.

Figure 15. Isobologram showing potentiating effect of the combination of taxol plus 10 Example B-3.

Figure 16. Isobologram showing potentiating effect of the combination of taxol plus Example B-4.

Figure 17. Isobologram showing potentiating effect of the combination of taxol plus Example B-5.

15 Figure 18. Isobologram showing potentiating effect of the combination of taxol plus Example B-6.

Figure 19. Isobologram showing potentiating effect of the combination of taxol plus Example B-7.

Figure 20. Isobologram showing potentiating effect of the combination of taxol plus 20 Example B-8.

Figure 21. Isobologram showing potentiating effect of the combination of taxol plus Example B-9.

Figure 22. Effect of KT5720 and taxol on the growth of MX-1 tumors.

Figure 23. Table of data showing toxicity of several of the drugs both individually 25 and in combination with taxol on non-tumored mice. (In Vivo Effects)

DETAILED DESCRIPTION OF THE INVENTION

The compounds of this invention are of two types. The first type are known compounds described here for their usefulness when combined with taxol type compounds and used to treat cancer. The second type of compounds are novel compounds described here for the first time. 30 These novel compounds are also useful when combined with taxol type compounds and used for the treatment of cancer.

I. Known Compounds

I. The known compounds, and the source of those compounds, are listed below, and described by name and by reference to the labeled structures.

35 A) Indolocarbazole Type Compounds

1) The "First Known Derivatives of K-252." The "First Known Derivatives

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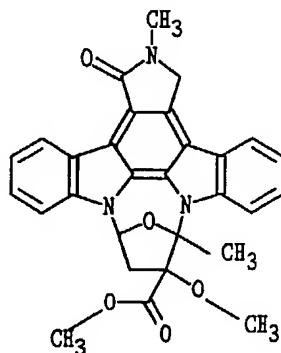
of K-252" are all of the compounds disclosed in U.S. patent 4, 877,776. U.S. patent 4, 877,776 incorporated herein by reference.

2) The "Second Known Derivatives of K-252." The "Second Known Derivatives of K-252" are all of the compounds disclosed in U.S. patent 4,923,986. U.S. patent 5 4,923,986 incorporated herein by reference.

3) The specific compounds below are more preferred,

3(a) KT5823

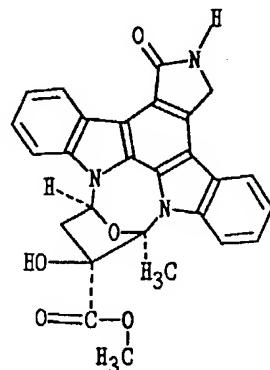
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3(b) K-252A

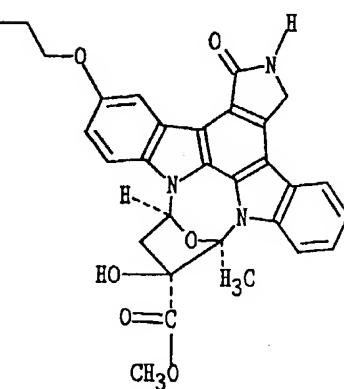
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3(c) KT5926

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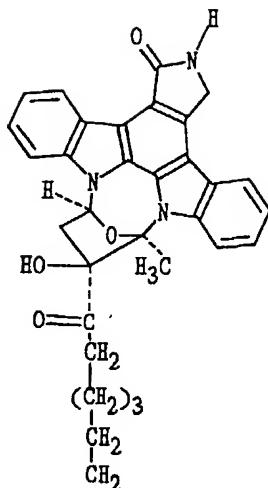
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3(d) KT5720

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3(e) Staurosporine

15 B. Non-Indolecarbazole Type Compounds

- 1) Adriamycin^s
- 2) Amiloride^s
- 3) Calphostin^s
- 4) Chlorpromazine^s
- 5) The compound known as "HA-1004"
- 6) Indomethacin^s
- 7) Okadaic acid
- 8) Phenazocine^s
- 9) Polymyxin B^s
- 10) 2-aminopurine^s
- 11) 6-dimethyl-aminopurine^s
- 12) Sphingosine^s
- 13) Tamoxifen
- 14) Compounds related to tamoxifen such as triphenylethylene antiestrogens
- 15) Trifluoperazine^s
- 16) Verapamil^s
- 17) 3-isobutyl-1-methyl-xanthine^s
- 18) 8-Cl-cAMP

35 Compounds marked with a superscript ^s are available from Sigma Chemical Company.
 Taxol and taxotere can be obtained from The National Cancer Institute.

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The clinical pharmacology of taxol is reviewed by Eric K. Rowinsky and Ross C. Donehower, The Clinical Pharmacology and Use of Antimicrotubule Agents in Cancer Chemotherapeutics, Pharmac. Ther., Vol 52, pp 35-84, 1991. Clinical and preclinical studies with taxol are reviewed by William J. Slichenmyer and Daniel D. Von Hoff, Taxol: A New and

5 Effective Anti-cancer Drug, Anti-Cancer Drugs, Vol. 2, pp 519-530, 1991.

Taxol and analogs thereof are the subject of various patents including, for example, U.S. Patent Nos. 4,814,470; 4,857,653; 4,942,184; 4,924,011; 4,924,012; 4,960,790; 5,015,744; 5,157,049; 5,059,699; 5,136,060; 4,876,399 as well as PCT Publication No. WO 92/09589, European Patent Application 90305845.1 (Publication No. A2 0 400 971), 89400935.6

10 (Publication No. A1 0 366 841) and 90402333.0 (Publication No. 0 414 610 A1), 87401669.4 (A1 0 253 739), and PCT Publication Nos. WO 91/17977, WO 91/17976, WO 91/13066, WO 91/13053.

Rebeccamycin is described in: T. Kaneko and H. Wong, Tetrahedron Letters, Vol. 26, No. 34, pp 4015-4018 (1985).

15 The compounds known as K252a, K252b, KT5720, KT5823, KT5926, okadaic acid and staurosporine, are available from Kamiya Biomedical Company, Thousand Oaks, California.

The compounds known as "H-7," "H-9" and "HA-1004" (B4-B6) are available from Seikagaku America, Inc., St. Petersburg, Florida

20 Lavendustin (B8) is available from Gibco BRL.

The compound, Kampferol-7-neonesperidoside, is available from Apin Chemical Co., Abingden, Oxfordshire, United Kingdom.

All of the documents referred to above are incorporated by reference herein.

II. New Compounds

II. The new compounds of this invention are identified in two ways: by the descriptive name and by reference to structures contained in appropriate charts. In some situations, the proper stereochemistry is also represented in the charts.

25 In this document the parenthetical term (C_n-C_m) is inclusive such that a compound of (C_1-C_8) would include compounds of one to 8 carbons and their isomeric forms. The various carbon moieties are defined as follows: Alkyl refers to an aliphatic hydrocarbon radical and

30 includes branched or unbranched forms such as methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, t-butyl, n-pentyl, isopentyl, n-hexyl, isohexyl, n-heptyl, isoheptyl, and n-octyl.

35 Alkoxy as represented by $-O-(C_1-C_8\text{ alkyl})$ refers to an alkyl radical which is attached to the remainder of the molecule by oxygen and includes branched or unbranched forms such as methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, sec-butoxy, t-butoxy, n-pentoxy, isopentoxy, n-hexaoxy, isohexaoxy, n-heptoxy, isoheptoxy, and n-octoxy.

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(C₃-C₁₀)cycloalkyl refers to a radical of a saturated cyclic hydrocarbon which includes alkyl-substituted cycloalkyl, such as cyclopropyl, 2-methylcyclopropyl, 2,2-dimethylcyclopropyl, 2,3 diethylcyclopropyl, 2-butylcyclopropyl, cyclobutyl, 2-methylcyclobutyl, 3-propylcyclobutyl, cyclopentyl, 2,2-dimethylcyclopentyl, cyclohexyl, cycloheptyl, or cyclooctyl. Each of these

5 moieties may be substituted as appropriate.

It will be apparent to those skilled in the art that compounds of this invention may contain chiral centers. The scope of this invention includes all enantiomeric or diastereomeric forms of formula I compounds either in pure form or as mixtures of enantiomers or diastereomers. The therapeutic properties of the compounds may to a greater or lesser degree
10 depend on the stereochemistry of a particular compound.

Both organic and inorganic acids can be employed to form non-toxic pharmaceutically acceptable acid addition salts of the compounds of this invention. Illustrative acids are sulfuric, nitric, phosphoric, hydrochloric, citric, acetic, lactic, tartaric, palmoic, methanesulfonic, ethanesulfonic, sulfamic, succinic, cyclohexylsulfamic, fumaric, maleic, and benzoic acid.

15 These salts are readily prepared by methods known in the art.

The compounds of this invention can be made in accordance with the processes described in the PREPARATIONS AND EXAMPLES for the preparation of novel compounds and illustrated in the GENERAL REACTIONS and the REACTIONS OF CHART A and CHART B.

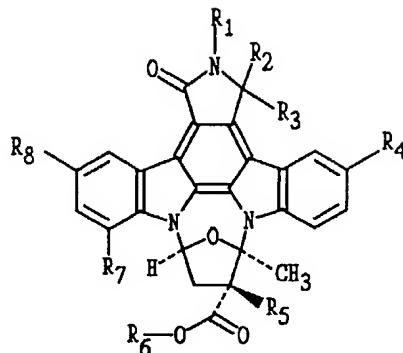
20 In clinical practice the compounds of the present invention will normally be administered by injection, in the form of pharmaceutical preparations comprising the active ingredient either as a free base or as a pharmaceutically acceptable non-toxic, acid addition salt, such as the hydrochloride, lactate, acetate, mesylate, methanesulfonate, or sulfamate salt, in association with a pharmaceutically acceptable carrier. The use and administration to a patient
25 to be treated in the clinic would be readily apparent to a physician or pharmacist of ordinary skill in the art.

In therapeutical treatment the suitable daily doses of the compounds of the invention should fall within the following ranges: Taxol, taxotere and related compounds should be administered from .001 mg/kg to 10 mg/kg, preferably between .05 mg/kg to 5 mg/kg for
30 intravenous administration. The compounds to be combined with taxol should be administered in the same dosage range. The precise dosage will be apparent to an ordinarily skilled physician or pharmacologist taking into account factors such as the age, weight, sex, and medical condition of the patient being treated. Also relevant is the potency of the particular compound and its ability to potentiate the effects of taxol. The potency of the compounds are indicated by
35 the standard tests described below.

The New Compounds:

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A compound of FORMULA I, below,



FORMULA I

wherein,

R₁ is -H, -(C₁-C₄ alkyl), -C(O)-(C₁-C₄ alkyl), -NH₂,
 -C(O)-NH₂, -CH₂CH₂-N(R₁₋₁)₂,

wherein R₁₋₁ is -H or -(C₁-C₄ alkyl),

15 R₂ is -H, or R₂ and R₃ taken together are (O),

R₃ is -H, -OH or R₂ and R₃ taken together are (O),

R₄ is -H, -OH, -NH₂, or -O-(C₁-C₄ alkyl),

R₅ is -OH, -O-(C₁-C₄ alkyl), or -O-C(O)-(C₁-C₄ alkyl),

R₆ is -(C₆-C₁₂ alkyl), -(C₃-C₁₀ cycloalkyl), -(CH₂)_nCH₂N(R₆₋₁)₂,

20 wherein R₆₋₁ is -H, or -(C₁-C₄ alkyl),

R₇ is -H, or -NH₂,

R₈ is -Cl, -Br, -H, -CH₃, -CH₂OH, -OH, -O-(C₁-C₄ alkyl),

-N(R₈₋₁)₂, or -NHC(O)-NH(R₈₋₁),

wherein R₈₋₁ is -H or -(C₁-C₄ alkyl)

25 wherein n is 0-5

with the proviso that:

a) when R₂ or R₃ is -OH then R₁ is H,

b) when R₁, R₂, R₃, R₄, and R₇ all equal H and R₅ is OH

then R₆ does NOT equal -(CH₂)₅CH₃,

30 c) when R₁, R₄, and R₇ all equal H, and R₂ combined with R₃ is (O),

and R₅ is OH, then R₆ does NOT equal -(CH₂)₅CH₃.

d) when R₄ is -OH, -NH₂, or -O-(C₁-C₄ alkyl), then R₄ and R₈ are the same.

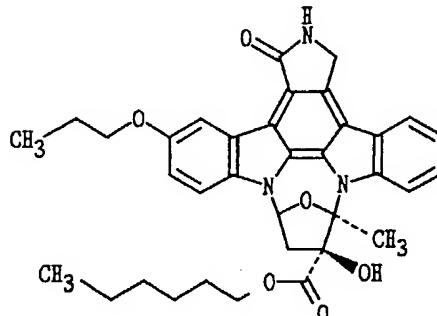
Preferred Compounds

35 The preferred compounds of this invention are those, referring to the compound of FORMULA I, wherein R₁ is H or CH₃; R₂, R₃, and R₇ is H; R₅ is OH or OCH₃;

R₈ is -O-(C₁-C₄ alkyl). The following compounds are preferred, Example B-4 and Example A-1, Example A-1 whose structure is shown below.

5

10



(Example A-1)

Biological Activity

Since taxol is known to be an effective chemotherapeutic agent, for example in the treatment of ovarian cancer, any co-treatment that increases the toxicity of taxol on cancer cells, such as CHO cells, would be likely to increase the chemotherapeutic effect of taxol in cancer patients or to allow smaller doses of taxol to be administered. The compounds of this invention synergize with taxol to produce tumor cell toxicity at lower doses than taxol alone, this requires the conclusion that the compounds will be effective in synergizing with taxol in killing tumor cells in human cancer patients. Additional studies that evaluate the compounds effects on human breast call MX-1 tumors, described below also support this conclusion.

The compounds of this invention are therefore useful for the same cancers for which taxol has been shown active, including human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia.

See, e.g., the clinical pharmacology of taxol is reviewed by Eric K. Rowinsky and Ross C. Donehower, *The Clinical Pharmacology and Use of Antimicrotubule Agents in Cancer Chemotherapy*, *Pharmac. Ther.*, Vol 52, pp 35-84, 1991. Clinical and preclinical studies with taxol are reviewed by William J. Slichenmyer and Daniel D. Von Hoff, *Taxol: A New and Effective Anti-cancer Drug*, *Anti-Cancer Drugs*, Vol. 2, pp 519-530, 1991.

Cell lines and growth.

The parental CHO line, 10001a, is a subclone of the CHO line Pro⁻⁵ (Stanley et al., 1975). The line was maintained in alpha-MEM Earle's Salts supplemented with 2 mM glutamine, 100 units/ml penicillin, 100 µg/ml streptomycin and 10% fetal bovine serum. All cell lines were maintained at 37°C in 5% CO₂ in a humidified incubator. Periodically, the cell lines were tested for mycoplasma and always found to be free of infection. Compounds were dissolved in dimethylsulfoxide (DMSO) and then diluted into medium for cell

growth assays.

Drug synergy experiments - "10001a" cell lines.

Cells were treated simultaneously with the experimental compound and taxol in 132 different combinations of doses in 96 well plates. The 96-well plates were incubated for four days. Cell growth was determined by the development of the colorimetric dye 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) as described by Mosmann, 1983. MTT dissolved in PBS at 2 mg/ml was added to the plates already containing growth medium to give a final concentration of 0.2 mg/ml in each well. Plates were then incubated for 3 hours. The medium containing MTT \pm drug was then aspirated off and 100 μ l/well isopropanol acidified with 0.04 N HCl was added. Plates were shaken for 5 minutes and absorbance was read at 570 nm on a Bio-tek EL 312e Bio-kinetics microplate reader.

Percent growth for 10001a cells was determined by dividing the absorbance reading at each drug dilution by the reading in control wells. LD₅₀s for each compound were determined to be the concentration of drug at which a 50% inhibition in cell growth was obtained.

15 Potentiating effects from the combination of compounds on 10001a cells was determined by graphing the combinations of drugs which gave LD₅₀s in the form of an isobologram (Kallman, 1987 and Brunden, 1988).

The effectiveness of combinations of compounds with taxol on the killing of wild type, 10001a, cells is shown by means of an isobologram. The compounds of this invention act in a potentiating or synergistic manner with taxol to kill cells with much lower doses in combination than would be expected if the drugs were merely exhibiting additive effects. This effect is surprising and unexpected. The isobolograms are displayed as FIGURES 1-21. FIGURES 1-21 demonstrate the effectiveness of combinations of compounds with taxol on the killing of wild type cells. FIGURE 4 is included in the series to show how a compound with no potentiating effect behaves.

The isobologram shows the effectiveness of a combination of 2 drugs for the killing of wild type, 10001a CHO, cells. The data line is the solid line with open circle or triangle data points. The data line shows the combination of doses which gives an LD₅₀ for the cells. In FIGURES 1-21 the concentration of taxol is plotted against the concentration of drug. The 30 diagonal dashed line shows the predicted concentrations of drugs if their combination only had an additive effect. If any data points were above the dashed line the date would indicate the combination of compounds had antagonistic effects. Data points below the line indicate the compounds have potentiating or synergistic effects. Compare the isobologram in FIGURE 4, showing NO potentiating effect, to the other isobolograms.

35 In addition to the data provided in the isobolograms the compounds have been tested in mice. Compound KT5720 has been tested on tumored mice and compounds KT5926 and

KT5720 have been tested in non-tumored mice.

Drug synergy experiments - MX-1 tumors.

FIGURE 22 shows the effect of KT5720 both separately and in combination with taxol in tumored mice. Human breast cell MX-1 tumors were implanted subcutaneously as 2 mm cubes in athymic mice. Mice were dosed every day for five days with drugs or vehicle control. The vehicle used was 2% dimethylacetamide, 10% emulphor, 88% saline. Animals received 12.5 mg/kg taxol (shown in figure as solid circle data points), 25 mg/kg KT5720 (shown in figure as open triangle data points), 12.5 mg/kg taxol + 25 mg/kg KT5720 (shown in figure as solid triangle data points), or vehicle alone (shown in figure as open square data points). Tumor burden was measured every two or three days starting with day 5 and volume was calculated. In FIGURE 22 the size of the tumor in millimeters is plotted against time in days. Eight mice were used per dose group. Results are graphed with standard errors. The results show that there was no effect of KT5720 alone on inhibition of growth of the tumor cells. Taxol, at 12.5 mg/kg, has a modest effect on reducing the tumor burden in these mice. The combination of KT5720 plus taxol clearly show a potentiation of the taxol effect by the addition of KT5720. In summary, KT5720 has no effect by itself, but in combination with taxol, at the dosage tested, it causes a dramatic inhibition of tumor growth.

Drug synergy experiments - non-tumored mice.

FIGURE 23 shows the effects of compounds KT5926 and KT5720 on non-tumored mice. When the compounds are combined with taxol and then administered to non-tumored normal mice they show a dramatic amount of toxicity. There was no lethality at the doses shown when these drugs were given individually. This means that there are strong synergistic effects with the compounds *in vivo*. The combination of drugs will be effective in tumor bearing mice and as a cancer treatment for humans. See FIGURE 23 - IN VIVO EFFECTS. This figure provides a table of data showing the toxicity of several of the drugs combined with taxol as compared to the individual administration of the drugs on non-tumored mice.

The required synergistically effective amounts (concentrations) will vary depending on the particular types of individual to be treated taking into consideration various conditions including age, weight, type of cancer treated, stage of disease, etc. Effective amounts can be readily determined by routine experimentation.

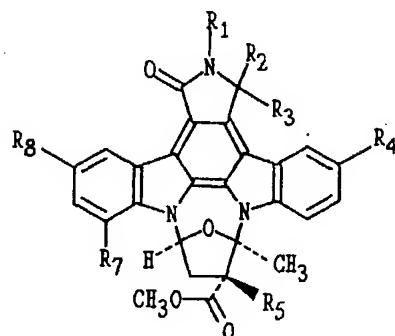
Without further elaboration, it is believed that one skilled in the art can, using the preceding description, practice the present invention to its fullest extent. The following detailed examples describe how to prepare the various compounds and/or perform the various processes of the invention and are to be construed as merely illustrative, and not limitations of the preceding disclosure in any way whatsoever. Those skilled in the art will promptly recognize appropriate variations from the procedures both as to reactants and as to reaction conditions and

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techniques.

PREPARATIONS AND EXAMPLES for the preparation of novel compounds.
GENERAL REACTIONS

5



10

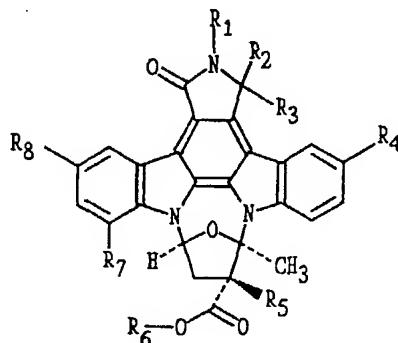
Starting materials

15

Step 1



20



25

Starting Materials. The starting materials are obtained by using the procedures described in U.S. Patent 4,923,986 and U.S. Patent 4,877,776. U.S. patents 4,923,986 and 4,877,776 are incorporated by reference into this document.

In addition to the patent references above, the starting materials for the reactions are also described in non-patent literature. The compound known as K252A is described in Kase, H., K. Iwahashi, and Y. Matsuda, K252a, "A Potent Inhibitor of Protein Kinase C from Microbial Origin." *J. Antibiot. (Tokyo)* 39:10066-1071 (1986). The compound known as KT 5926, and related compounds are described in S. Nakanishi, K. Yamada, K. Iwahashi, K. Kuroda and H. Kase, "KT5926, a Potent and Selective Inhibitor of Myosin Light Chain Kinase." *Molecular Pharmacology*, 37:482-488 (1990). The other compounds of formula 1 where R6 is C1-C5 alkyl are described in U.S. Patent 4,923,986 and U.S. patent 4,877,776. In general, treatment of

-17-

compounds of formula 1, where R_6 is CH_3 , are treated with R_6 -OH where (R_6 is C_6 - C_{12} alkyl) and KCN to give the desired compounds. All the starting materials are described in the above patents. Compounds of the type R_5 is H are described in WO 91/09034 published 27 June 1991. All the above documents are incorporated by reference herein.

5 The General Procedure for producing variations for the R_6 group is as follows:

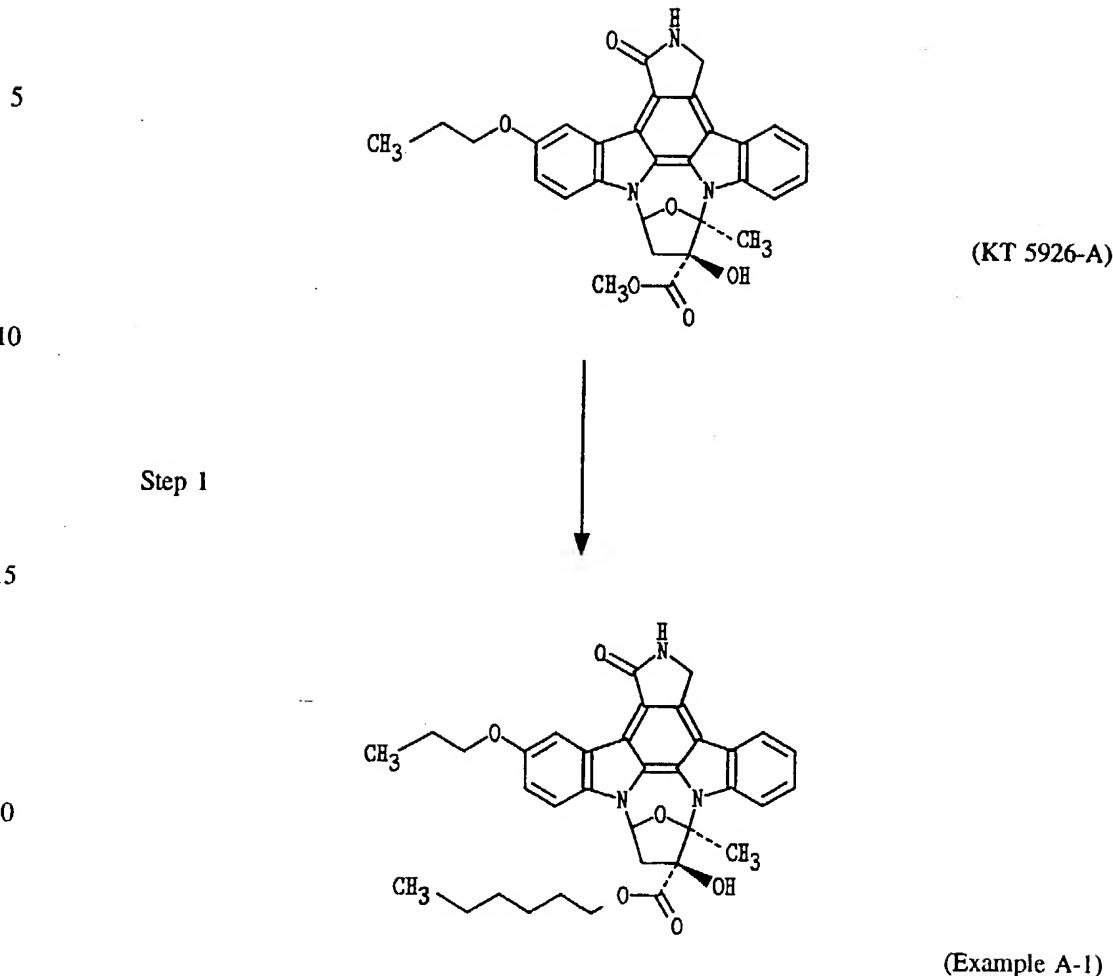
To an appropriate starting material such as KT252a add an alcohol such as n-hexanol (R_6 is $-(CH_2)_5CH_3$). Stir the mixture at temperatures ranging from room temperature to 125 degrees until dissolution is complete. An equal weight amount of solid KCN is added and the reaction mixture is stirred for an additional 18 to 144 hours at temperatures ranging from room

10 temperature to 125 degrees. The reaction mixture is poured into ethyl acetate and the ethyl acetate solution is extracted with water. The organic solution is dried over anhydrous sodium sulfate, filtered and evaporated to dryness under high vacuum at temperatures ranging from 35° to 70°C to near dryness. Hexane is added to the residue and the resulting solids are allowed to sit, under hexane, for 24 hours. The solids are filtered and washed well with hexane and dried

15 at 40°C to give the desired compound. The materials may be identified by their retention time on HPLC. Detailed HPLC conditions are provided in the examples below. For example, when the following HPLC conditions are used the retention time for KT252a is 2.54 minutes. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute).

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REACTIONS OF CHART A



25

Procedure A. Reactions of step 1, above. Preparation of **Example A-1**, (R_8 is $-O-(CH_2)_2CH_3$, R_6 is $-(CH_2)_5CH_3$), from the starting material KT-5926.

Example A-1 is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-30 10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9- methyl-1-oxo-16-propoxy-, hexyl ester, (9R-(9.alpha.,10.beta., 12.alpha.)).

To 0.8 mg of KT-5926 (0.0015 mmol) is added 0.5 ml of n-hexanol and, after dissolution is complete, 1.0 mg of KCN is added. The reaction mixture is stirred at room temperature for 24 hours at which time HPLC indicates all the starting material has reacted.

35 Add 0.5 ml of a 35:65 water:acetonitrile solution, followed by acetonitrile (to complete dissolution) and chromatograph the entire reaction solution on a preparatory HPLC system (2-

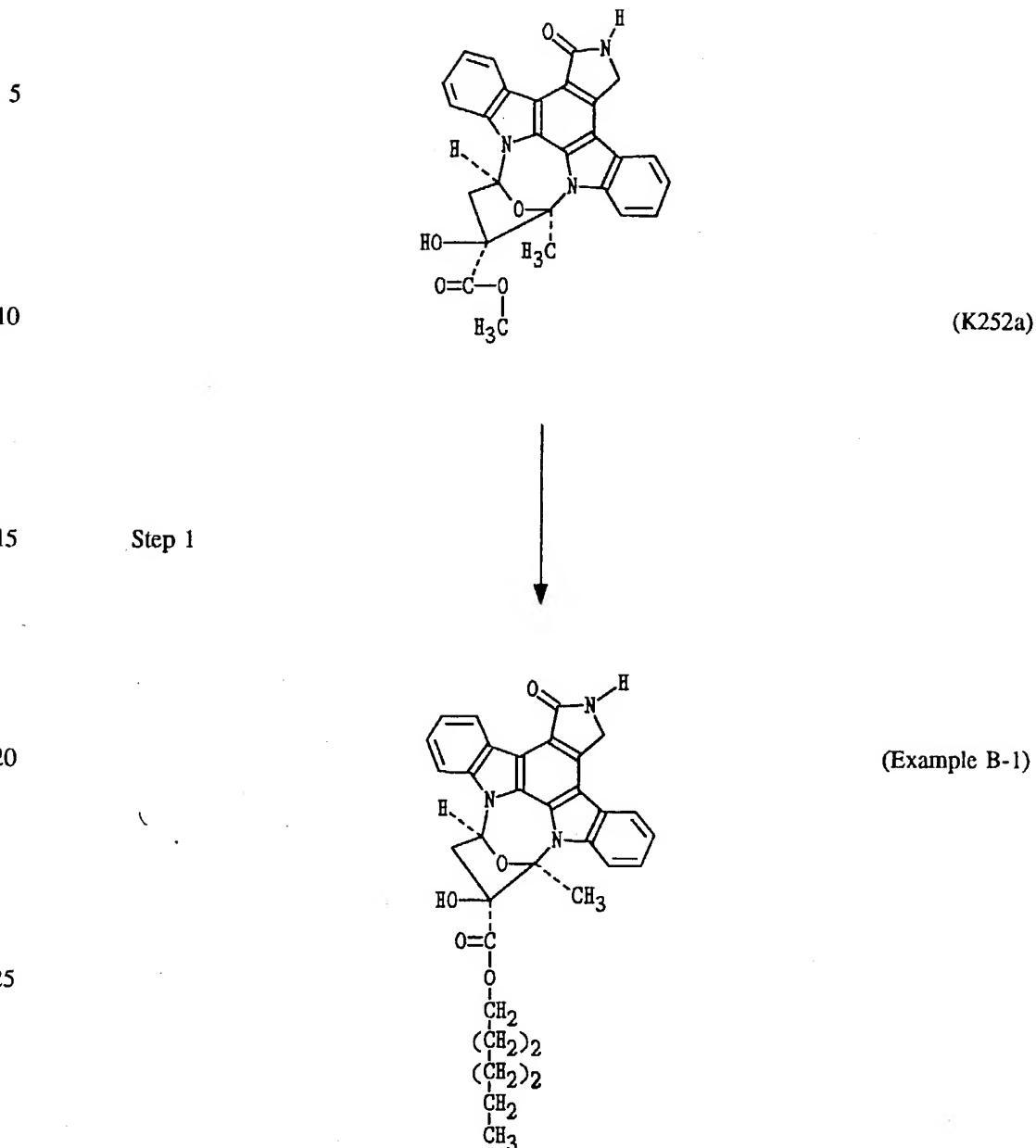
-19-

PrepPak 25 X 100 mm Cartridge) microBondapak C18, 10 microns; 35:65 water:acetonitrile at 8 ml/min taking 24 ml fractions. The fractions containing Example A-1, are combined and evaporated to dryness to give 0.12 mg of product. HPLC data was run as follows: HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute) rt is

5 18.69 minutes for Example A-1; rt is 3.63 minutes for KT 5926.

-20-

REACTIONS OF CHART B



30 Procedure B. Reactions of step 1, above. Preparation of Example B-1, (R_8 is H, R_6 is $-(CH_2)_6CH_3$), from the starting material, K252a.

Example B-1 is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, heptyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)).

To 15 mg of K252a (0.032 mmol) add 2 ml of n-heptanol (R_6 is

$-(\text{CH}_2)_6\text{CH}_3$. The mixture is allowed to stir at room temperature until dissolution is complete. 15 mg of KCN is added and the reaction mixture is stirred for an additional 96 hours. The reaction mixture is poured into 20 ml of ethyl acetate and the ethyl acetate solution is extracted with water. The organic solution is dried over anhydrous sodium sulfate, filtered and evaporated to dryness under high vacuum at 59°C to near dryness. 15 ml of hexane is added to the residue and the resulting solids are allowed to sit, under hexane, for 24 hours. The solids are filtered and washed well with hexane and dried at 40°C to give 7.8 mg of Example B-1. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute) retention time is 7.14 minutes for Example B-1; 2.54 minutes for K252a. Mass Spec. theory predicts 552.2498 mass units; Measured: 552.2484.

Example B-2, (R₆ is $-(\text{CH}_2)_7\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, octyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)).

Using procedure B only substituting n-octanol (R₆ is $-(\text{CH}_2)_7\text{CH}_3$) in the reaction described above, and stirring at room temperature for 120 hours, gives Example B-2, as an amber solid. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute) retention time is 3.325 minutes. Mass Spec. theory predicts 566.2655 mass units; Measured: 566.2652.

Example B-3, (R₆ is $-(\text{CH}_2)_8\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, nonyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)).

Using procedure B only substituting n-nonanol (R₆ is $-(\text{CH}_2)_8\text{CH}_3$) in the reaction described above, and stirring at room temperature for 144 hours, gives Example B-3, as an amber solid. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (25:75 water:acetonitrile; 2 ml/minute) retention time is 6.54 minutes. Mass Spec. theory predicts 580.2811 mass units; Measured: 580.2819

Example B-4, (R₆ is $-\text{CH}(\text{CH}_2\text{CH}_3)((\text{CH}_2)_3\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 1-ethylpentyl ester.

Using procedure B only substituting 3-heptanol (R₆ is $-\text{CH}(\text{CH}_2\text{CH}_3)((\text{CH}_2)_3\text{CH}_3$) in

the reaction described above, stirring at 110 degrees for 144 hours, gives Example B-4, as an amber solid. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute) retention time is 6.44 minutes. Mass Spec. theory predicts 552.2498 mass units; Measured: 552.2501.

5

Example B-5, (R_6 is $-\text{CHCH}_3(\text{CH}_2)_4\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-methylhexyl ester.

10 Using procedure B only substituting 2-heptanol (R_6 is $-\text{CHCH}_3(\text{CH}_2)_4\text{CH}_3$) in the reaction described above, and heating at 100 degrees for 18 hours, gives Example B-5, as an amber solid. HPLC retention time is 6.67 minutes. Mass Spec. theory predicts 552.2498 mass units; Measured: 552.2501.

15 **Example B-6**, (R_6 is $-\text{CHCH}_3(\text{CH}_2)_5\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-methylheptyl ester.

20 Using procedure B only substituting 2-octanol (R_6 is $-\text{CHCH}_3(\text{CH}_2)_5\text{CH}_3$) in the reaction described above, stirring at 100 degrees for 96 hours, gives Example B-6, as an amber solid. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (25:75 water:acetonitrile; 2 ml/minute) retention time is 4.55 minutes. Mass Spec. theory predicts 566.2655 mass units; Measured: 566.2652.

25 **Example B-7**, (R_6 is $-\text{CHCH}_3(\text{CH}_2)_6\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-methyloctyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)).

30 Using procedure B only substituting 2-nonanol (R_6 is $-\text{CHCH}_3(\text{CH}_2)_6\text{CH}_3$) in the reaction described above, stirring at 100 degrees for 96 hours, gives Example B-7, as an amber solid. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (25:75 water:acetonitrile; 2 ml/minute) retention time is 6.12 minutes. Mass Spec. theory predicts 580.2811 mass units; Measured: 580.2797.

35 **Example B-8**, (R_6 is $-(\text{CH}_2)_2\text{OCH}_2\text{CH}_3$), is named:

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-

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10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-,2-ethoxyethyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)).

Substituting ethoxyethanol (R_6 is $-(CH_2)_2OCH_2CH_3$) in the reaction described above, and stirring at room temperature for 96 hours, followed by 50 degrees for 24 hours, gives

5 Example B-8, as an amber solid. HPLC: Altex Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute) retention time is 1.54 minutes. Mass Spec. theory predicts 526.1978 mass units; Measured: 526.1959.

Example B-9, (R_6 is $-CH(cyclo-CH_2)_5$), is named:

10 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, cyclohexyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)).

Substituting cyclohexanol (R_6 is $-CH(cyclo-CH_2)_5$) in the reaction described above, stirring at 80 degrees for 18 hours, gives Example B-9, as an amber solid. HPLC: Altex 15 Ultrasphere-ODS 0.5 micron, 4.6 mm X 25 cm, (35:65 water:acetonitrile; 2 ml/minute) retention time is 3.95 minutes. Mass Spec. theory predicts 536.2185 mass units; Measured: 536.2169.

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TABLE 1 - IN VIVO EFFECTS

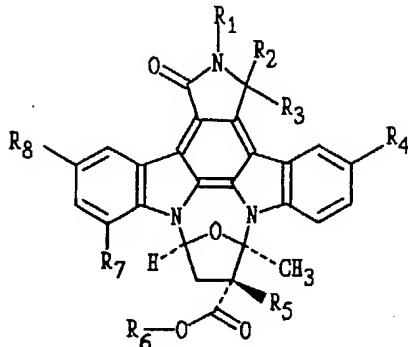
	mg/kg Dose	Day of Death	
		Mouse #1	Mouse #2
	Taxol	10.0	*--
		5.0	--
5	KT5926	15.0	--
		5.0	--
		1.0	--
		0.1	--
	Taxol + KT5926	10.0 + 15.0	10
10		10.0 + 5.0	7
		10.0 + 1.0	9
		10.0 + 0.1	9
		5.0 + 15.0	9
		5.0 + 5.0	14
15		5.0 + 1.0	10
		5.0 + 0.1	8
			9
20	BDF1 mice		
	Two mice per dose (mouse #1 and mouse #2)		
	Drugs were given intraperitoneally on days 1-9		
	* No day of death given means that the mice did not die during the duration of the experiment.		
25			

CLAIMS

1. A compound of FORMULA I, below,

5

10



FORMULA I

wherein,

R₁ is -H, -(C₁-C₄ alkyl), -C(O)-(C₁-C₄ alkyl), -NH₂,
-C(O)-NH₂, -CH₂CH₂-N(R₁₋₁)₂;

15 wherein R₁₋₁ is -H or -(C₁-C₄ alkyl);

R₂ is -H, or R₂ and R₃ taken together are, (O);

R₃ is -H, -OH or R₂ and R₃ taken together are, (O);

R₄ is -H, -OH, -NH₂, or -O-(C₁-C₄ alkyl);

R₅ is -OH, -O-(C₁-C₄ alkyl), or -O-C(O)-(C₁-C₄ alkyl);

20 R₆ is -(C₆-C₁₂ alkyl), -(C₃-C₁₀ cycloalkyl), (C₁-C₅ alkyl)-O-(C₁-C₅ alkyl),
-(CH₂)_nCH₂N(R₆₋₁)₂;

wherein R₆₋₁ is -H, or -(C₁-C₄ alkyl);

R₇ is -H, or -NH₂;

R₈ is -Cl, -Br, -H, -CH₃, -CH₂OH, -OH, -O-(C₁-C₄ alkyl);

25 -N(R₈₋₁)₂, or -NHC(O)-N(R₈₋₁)₂;

wherein R₈₋₁ is -H or -(C₁-C₄ alkyl);

wherein n is 0-5

with the proviso that:

a) when R₂ or R₃ is -OH then R₁ is H;

30 b) when R₁, R₂, R₃, R₄, and R₇ all equal H and R₅ is OH

then R₆ does NOT equal -(CH₂)₅CH₃;

c) when R₁, R₄, and R₇ all equal H, and R₂ combined with R₃ is (O)
and R₅ is OH, then R₆ does NOT equal -(CH₂)₅CH₃;

d) when R₄ is -OH, -NH₂, or -O-(C₁-C₄ alkyl), then R₄ and R₈ are the same.

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2. A compound of claim 1 wherein

R₁ is -H, -(C₁-C₄ alkyl), -C(O)-(C₁-C₄ alkyl), -C(O)-NH₂.

3. A compound of claim 1 wherein

5 R₁ is H or -CH₃.

4. A compound of claim 1 wherein

R₁ is -CH₂CH₂-N(R₁₋₁)₂.

10 5. A compound of claim 1 wherein

R₂ and R₃ are H.

6. A compound of claim 1 wherein

R₅ is -OH.

15

7. A compound of claim 1 wherein

R₅ is -OCH₃.

8. A compound of claim 1 wherein

20 R₆ is -(C₇-C₁₂ alkyl), -(C₃-C₁₀ cycloalkyl), -(CH₂)_nCH₂N(R₆₋₁)₂.

9. A compound of claim 1 wherein

R₆ is -(C₈-C₁₂ alkyl), -(C₃-C₁₀ cycloalkyl), -(CH₂)_nCH₂N(R₆₋₁)₂.

25 10. A compound of claim 2 wherein

R₂ and R₃ is (O).

11. A compound of claim 2 wherein

R₁ is -H, -(C₁-C₄ alkyl);

30 R₂ and R₃ are H and

R₅ is -OH or -O-(C₁-C₄ alkyl).

12. A compound of claim 5 wherein

R₅ is -OH or -O-(C₁-C₄ alkyl).

13. A compound of claim 5 wherein

R₈ is -Cl, -Br, -H, -CH₃, -CH₂OH, -OH, -O-(C₁-C₄ alkyl),
-N(R₈₋₁)₂, or -NHC(O)-NHR₈₋₁.

5 14. A compound of claim 5 wherein

R₈ is -O-(C₁-C₄ alkyl).

15. A compound of claim 11 wherein

R₁ is -H;

10 R₈ is -O-(C₁-C₄ alkyl).

16. A compound of claim 15 which is the compound named,

9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-
10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-16-propoxy-, hexyl
15 ester, (9R-(9.alpha.,10.beta., 12.alpha.)). (Example A-1)

17. A compound of claim 11 selected from the following named compounds,

- a) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-
10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, heptyl ester,
20 (9R-(9.alpha.,10.beta.,12.alpha.)), (Example B-1)
- b) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-
10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 1-ethylpentyl ester,
(Example B-4) or
- c) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-
25 10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-methylhexyl ester
(Example B-5).

18. A compound of claim 11 wherein

R₁ is -H;

30 R₆ is -(C₈-C₁₂ alkyl) and

R₈ is -O-(C₁-C₄ alkyl).

19. A compound of claim 11 wherein

R₁ is -H;

R₆ is -(C₃-C₁₀ cycloalkyl) or (C₁-C₅ alkyl)-O-(C₁-C₅ alkyl) and

R₈ is -O-(C₁-C₄ alkyl).

5

20. A compound of claim 11 wherein

R₁ is -H;

R₆ is -(C₈-C₁₂ alkyl) and

R₈ is H.

10

21. A compound of claim 20 selected from the following named compounds,

a) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, octyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)); (Example B-2)

15

b) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, nonyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)); (Example B-3)

c) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-methylheptyl ester,

20 (Example B-6)

d) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-methyloctyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)) (Example B-7).

25 22. A compound of claim 11 wherein

R₁ is -H;

R₆ is -(C₃-C₁₀ cycloalkyl) or (C₁-C₅ alkyl)-O-(C₁-C₅ alkyl) and

R₈ is H.

30 23. A compound of claim 22 selected from,

a) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, 2-ethoxyethyl ester, (9R-(9.alpha.,10.beta.,12.alpha.)); (Example B-8) or

b) 9,12-Epoxy-1H-diindolo(1,2,3-fg:3',2',1'-kl)pyrrolo(3,4-i)(1,6)benzodiazocine-

35 10-carboxylic acid, 2,3,9,10,11,12-hexahydro-10-hydroxy-9-methyl-1-oxo-, cyclohexyl ester,

(9R-(9.alpha.,10.beta.,12.alpha.)) (Example B-9).

24. A pharmaceutical composition for treating, controlling or preventing cancerous growths, such as human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia in mammals and humans, which comprises administering a therapeutic or prophylactic dosage of a compound of FORMULA I in conjunction with an appropriate dose of taxol or taxol related compounds.

5

25. A pharmaceutical composition for treating, controlling or preventing cancerous growths, such as human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia, in mammals and humans, which comprises administering a therapeutic or prophylactic dosage of any one of the compounds selected from the list below, in conjunction with an appropriate dose of taxol or taxol related compounds, a) KT5823, b) K252a; c) KT5926, d) KT5720, e) Staurosporine, f) drimycin, b) Amilorides, c) Calphostin, d) Chlorpromazine, e) The compound known as "HA-1004", f) Indomethacin, g) Okadaic acid, h) Phenazocine, i) Polymyxin, j) 2-aminopurine k) 6-dimethylaminopurine, l) Sphingosine, m) Tamoxifen, n) Compounds related to tamoxifen such as triphenylethylene antiestrogens, o) Trifluoperazine, p) Verapamil, q) 3-isobutyl-1-methyl-xanthine, r) 8-Cl-cAMP, in conjunction with an appropriate dose of taxol or taxol related compounds.

10

15

20 26. Use of a therapeutic or prophylactic dosage of a compound of FORMULA I in conjunction with an appropriate dose of taxol or taxol related compounds for the manufacture of a medicament for the treatment, control or prevention of cancerous growths, such as human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia in mammals and humans.

25

27. Use of a therapeutic or prophylactic dosage of the compounds described in U.S. Patents 4,877,776 or 4,923,986, in conjunction with an appropriate dose of taxol or taxol related compounds for the manufacture of a medicament for the treatment, control or prevention of cancerous growths, such as human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia in mammals and humans.

30

-30-

28. Use of a therapeutic or prophylactic dosage of any of the compounds selected from the following: a) KT5823, b) K252a, c) KT5926, d) KT5720, or e) Staurosporine, in conjunction with an appropriate dose of taxol or taxol related compound for the manufacture of a medicament for the treatment, control or prevention of cancerous growths, such as human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia in mammals and humans.

29. Use of a therapeutic or prophylactic dosage of any of the compounds selected from the following: a) Adriamycin
10 b) Amilorides
c) Calphostin
d) Chlorpromazine
e) The compound known as "HA-1004"
f) Indomethacin
15 g) Okadaic acid
h) Phenazocine
i) Polymyxin B
j) 2-aminopurine
k) 6-dimethyl-aminopurine
20 l) Sphingosine
m) Tamoxifen
n) Compounds related to tamoxifen such as triphenylethylene antiestrogens
o) Trifluoperazine
p) Verapamil
25 q) 3-isobutyl-1-methyl-xanthine or
r) 8-Cl-cAMP

in conjunction with an appropriate dose of taxol or taxol related compound for the manufacture of a medicament for the treatment, control or prevention of cancerous growths, such as human ovarian tumors, mammary tumors, and malignant melanoma, lung tumors, gastric tumors, colon tumors, head and neck tumors, and leukemia in mammals and humans.

FIGURE 1

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FIGURE 1.

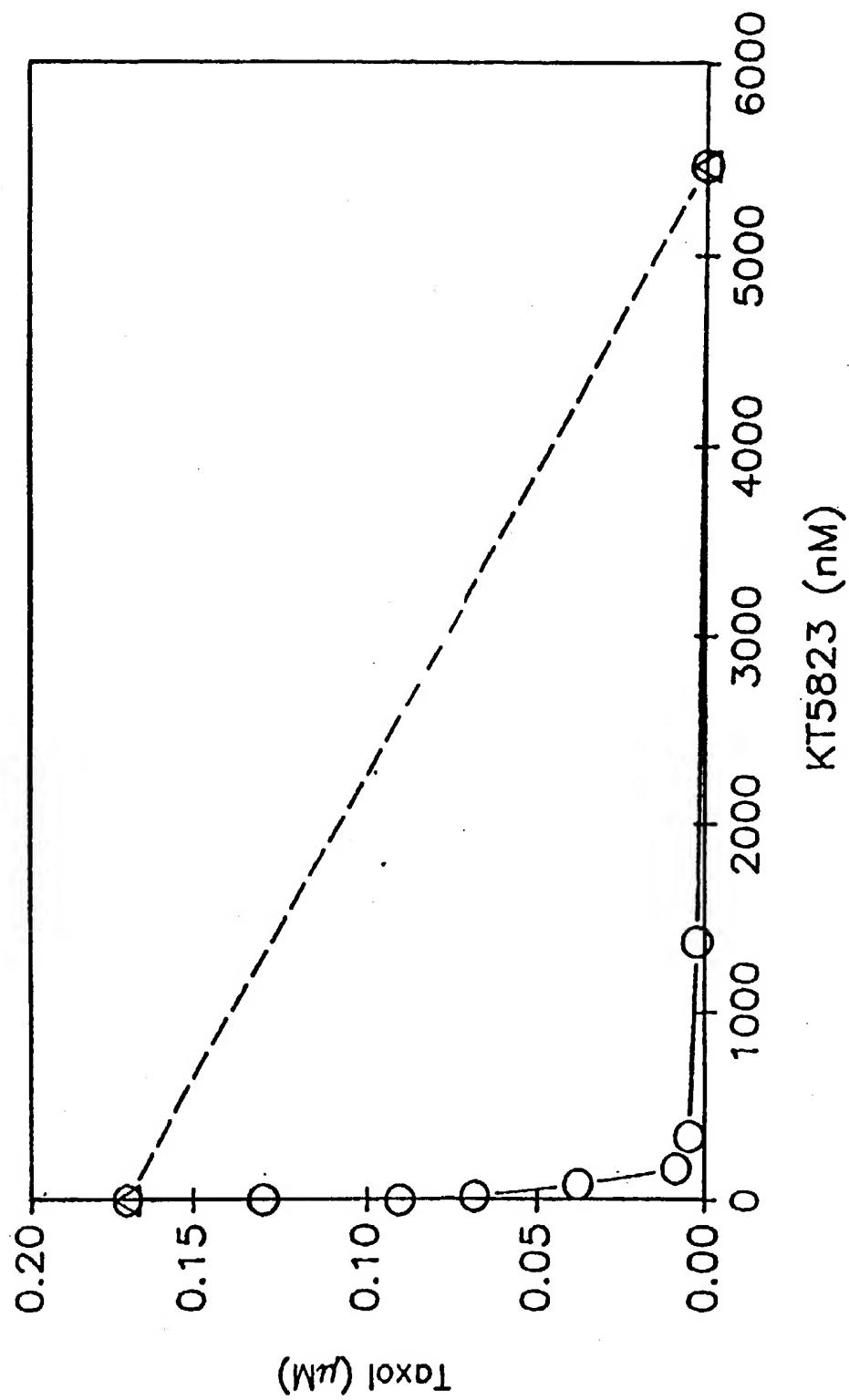
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FIGURE 2

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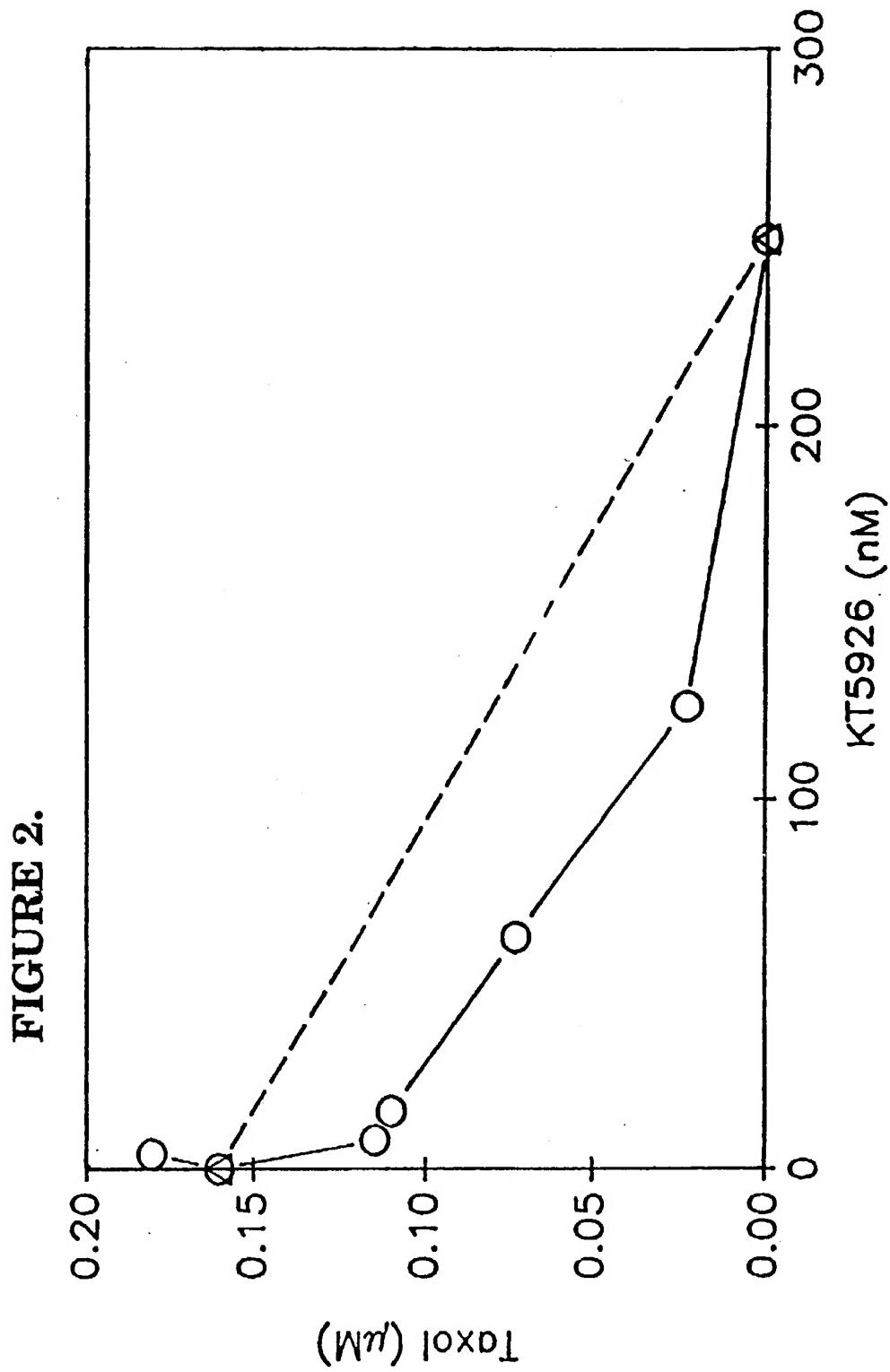
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FIGURE 3

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FIGURE 3.

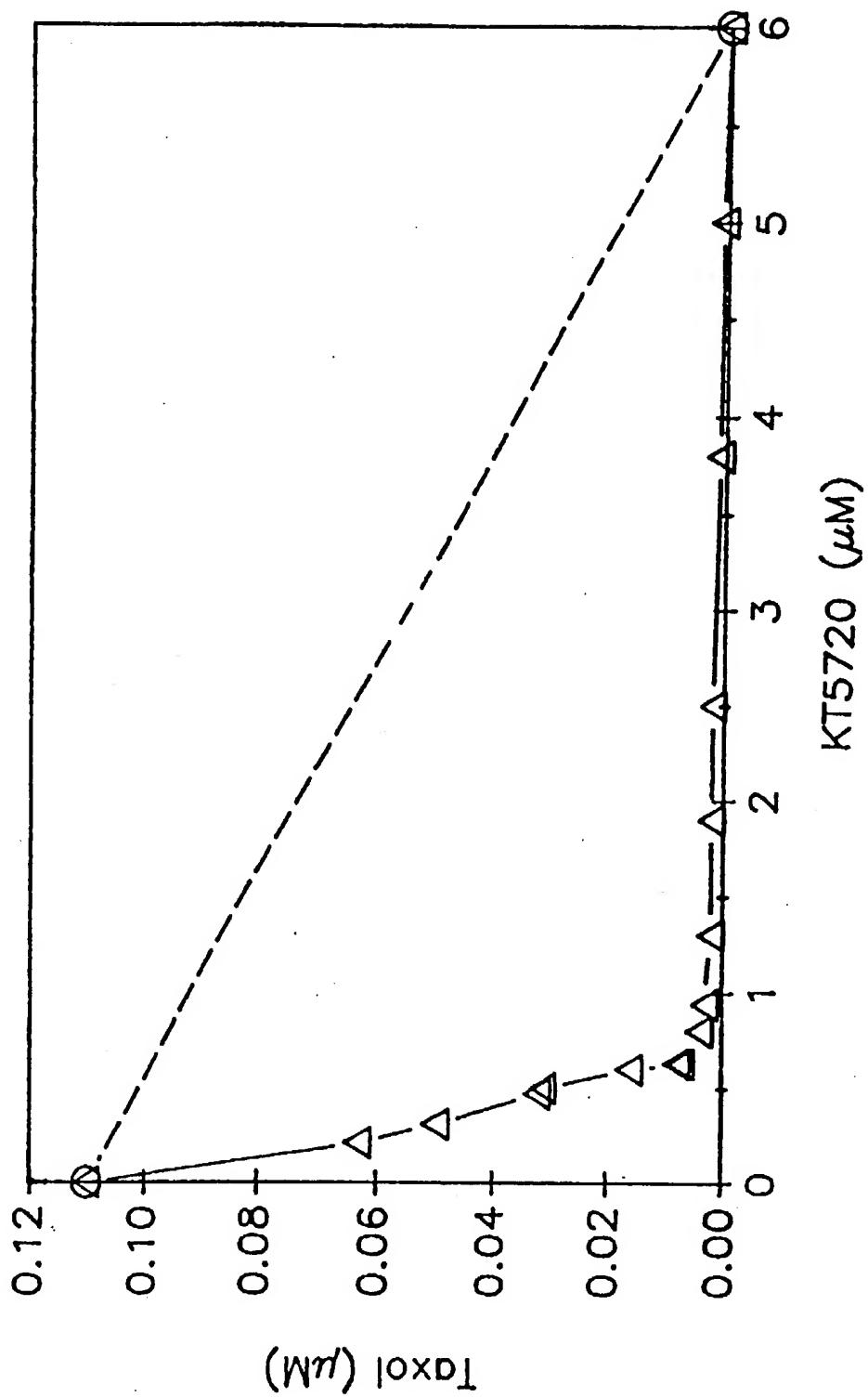
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FIGURE 4

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FIGURE 4.

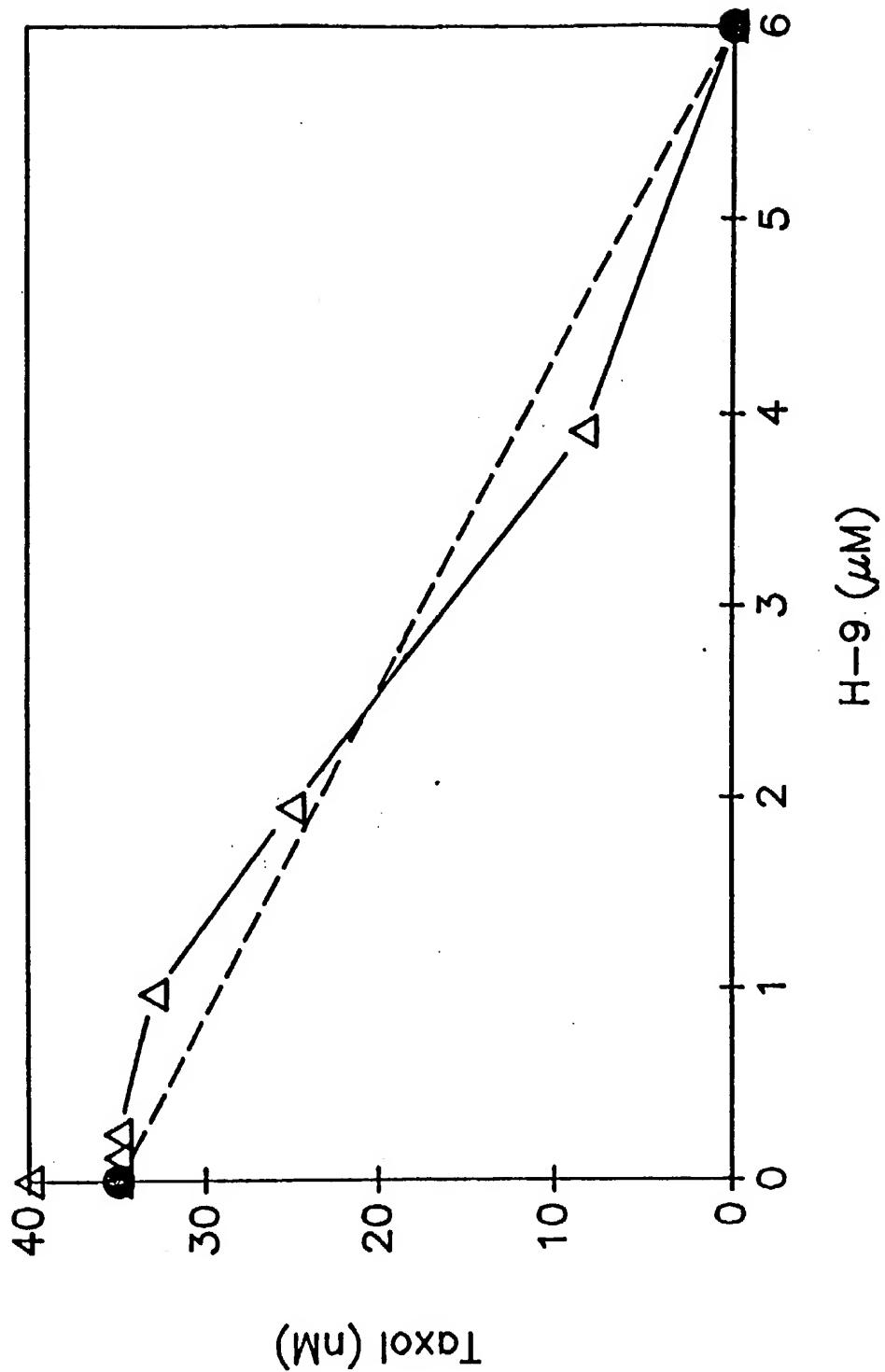


FIGURE 5

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FIGURE 5.

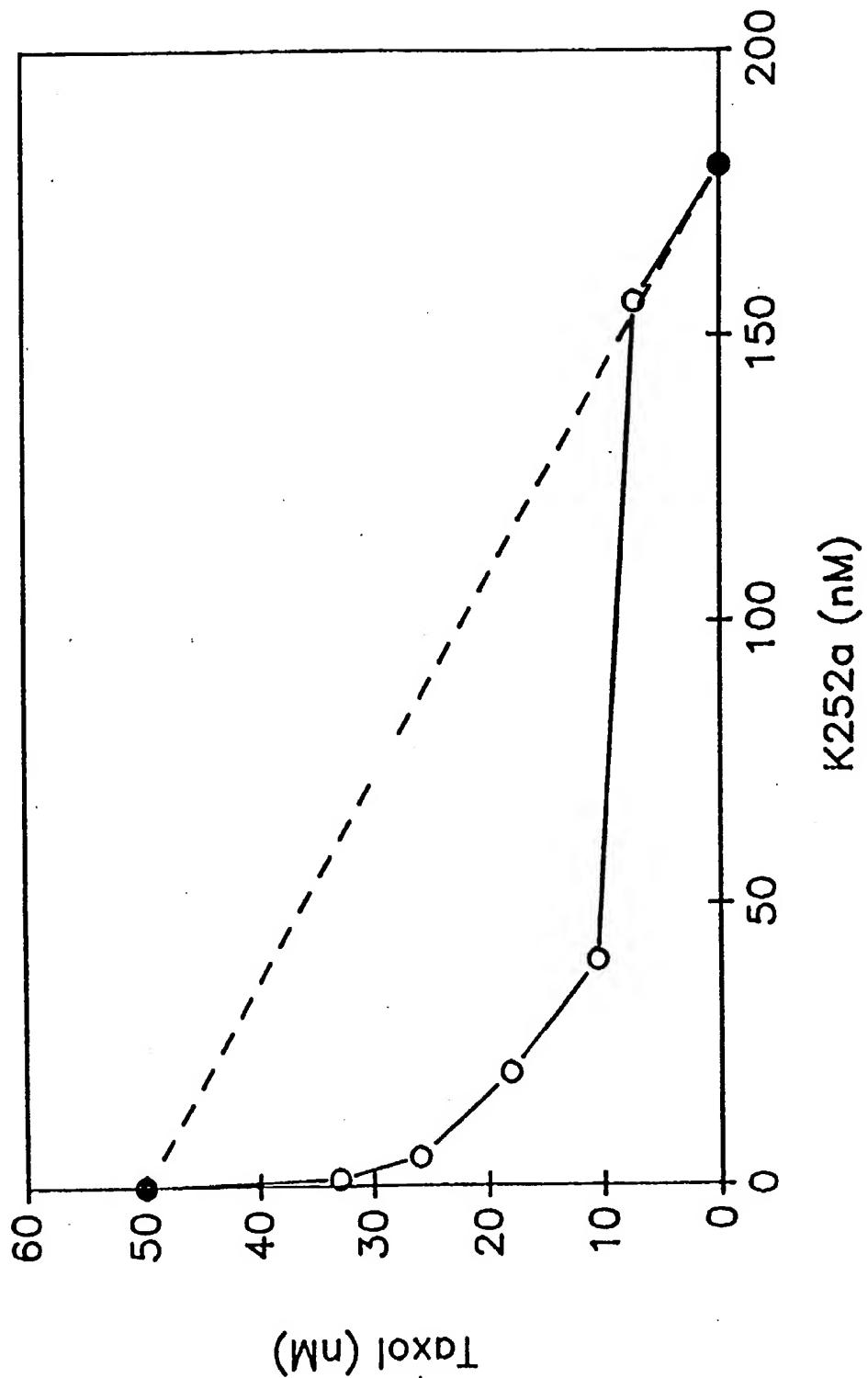


FIGURE 6.

FIGURE 6

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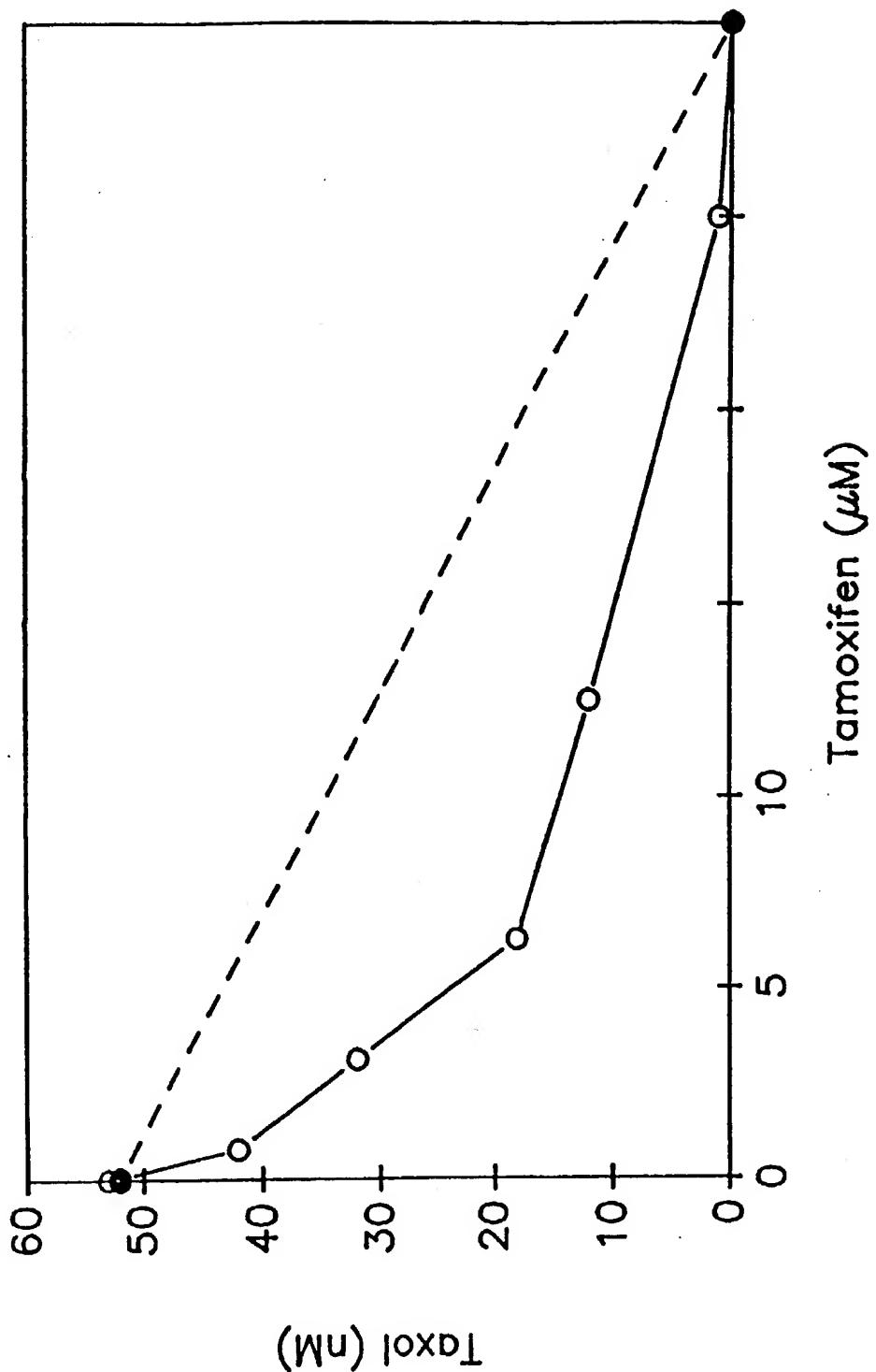


FIGURE 7

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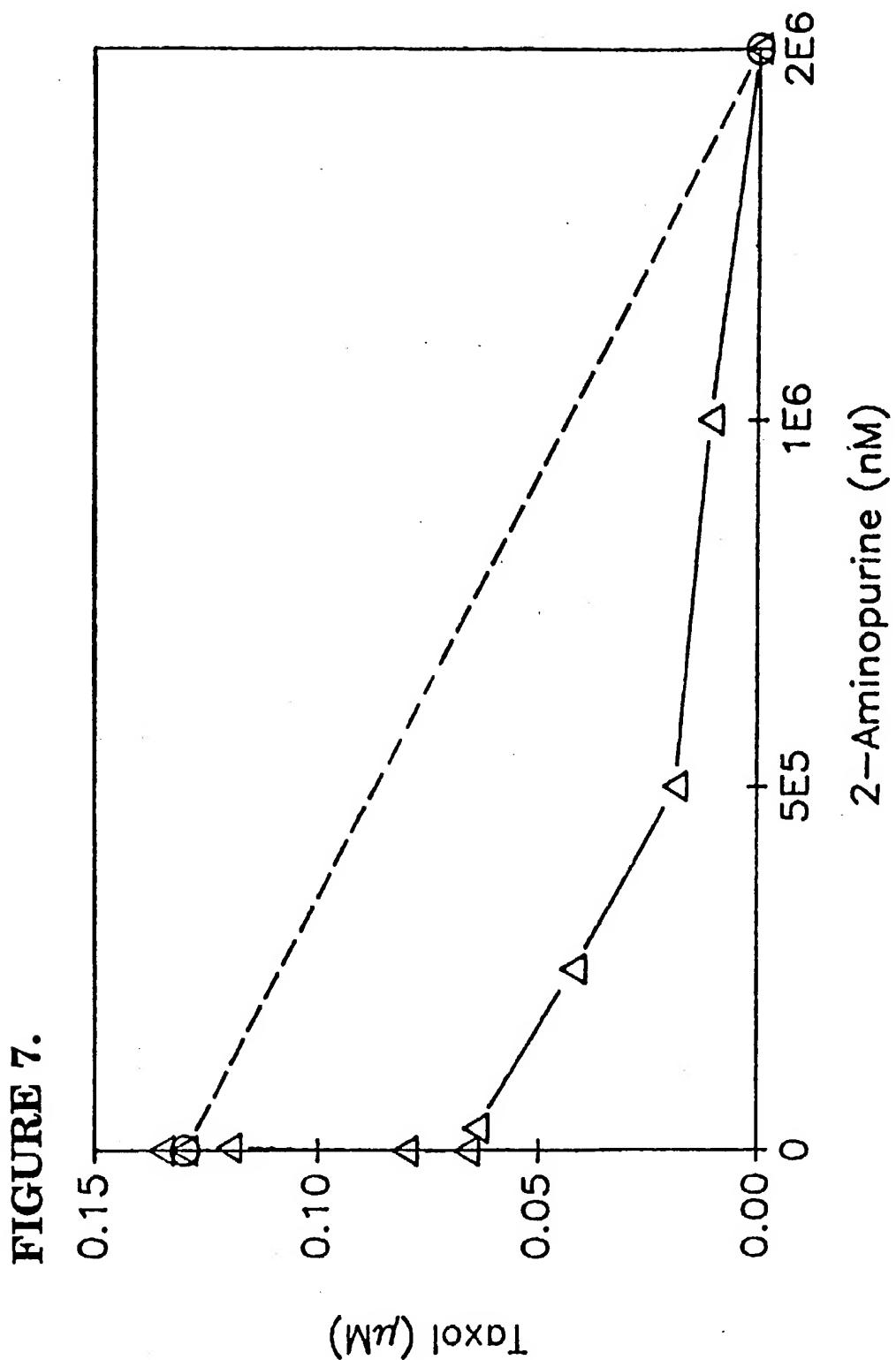


FIGURE 7.

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FIGURE 8

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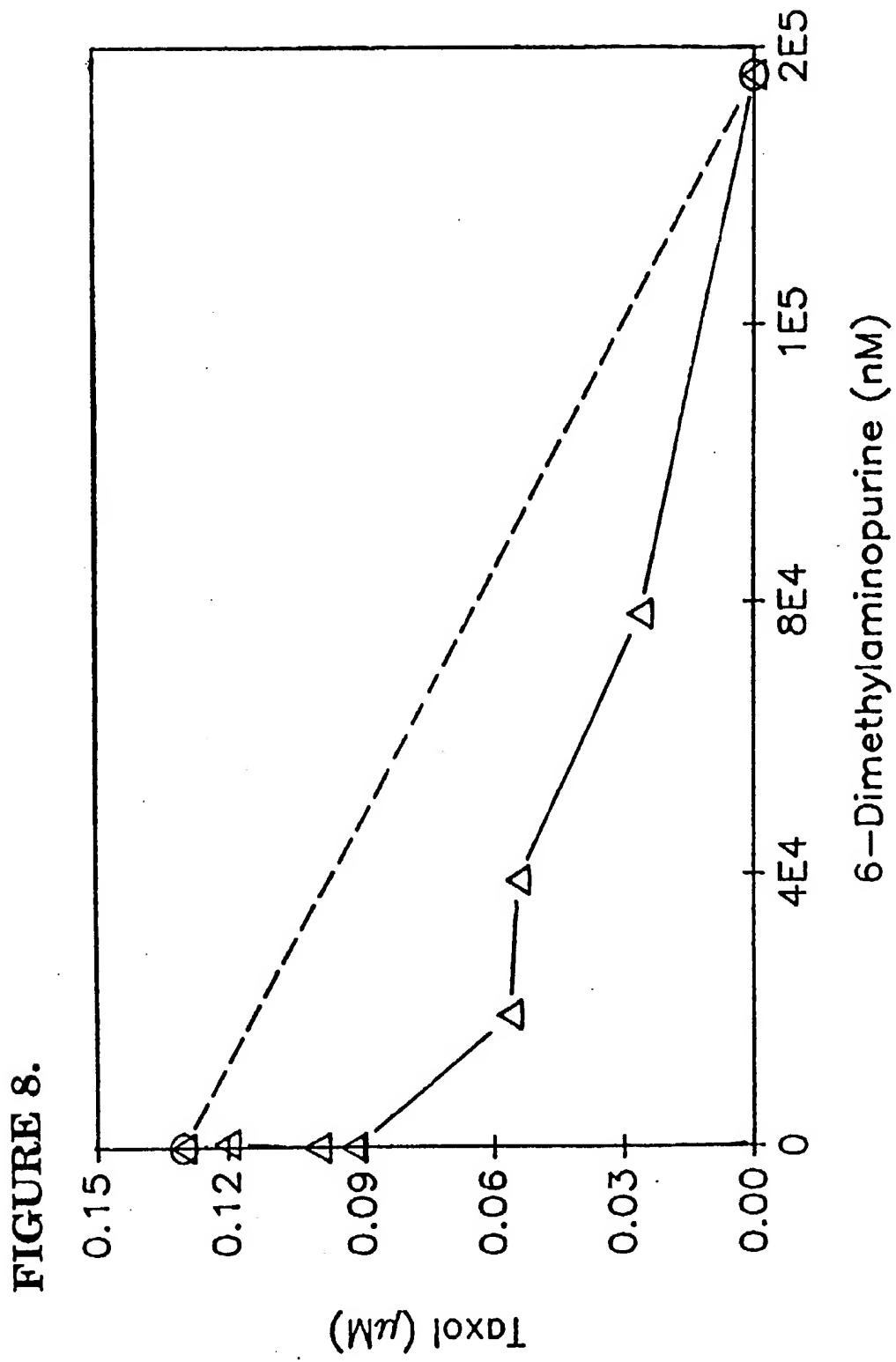
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FIGURE 9

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FIGURE 9.

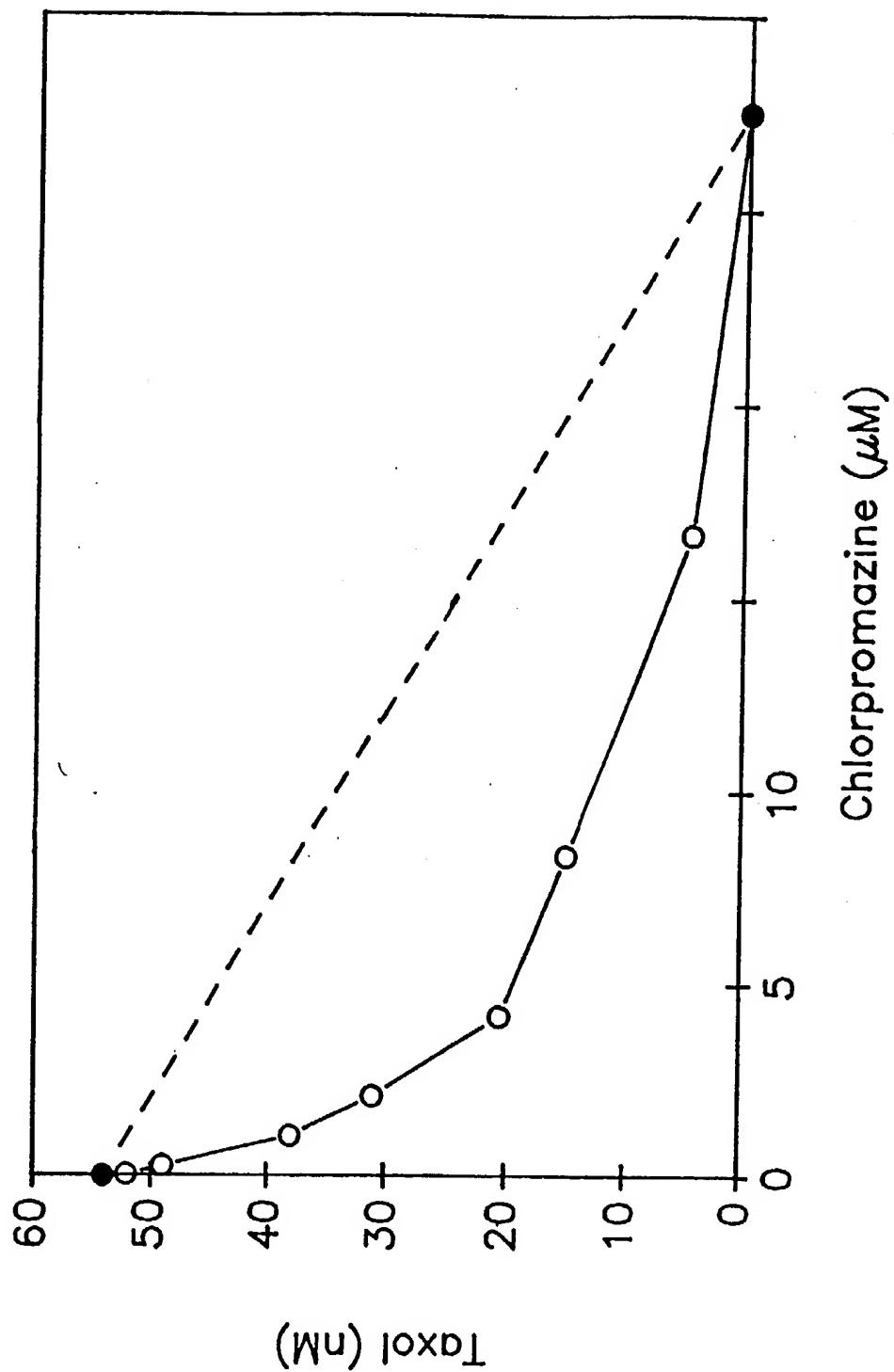
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FIGURE 10.

FIGURE 10

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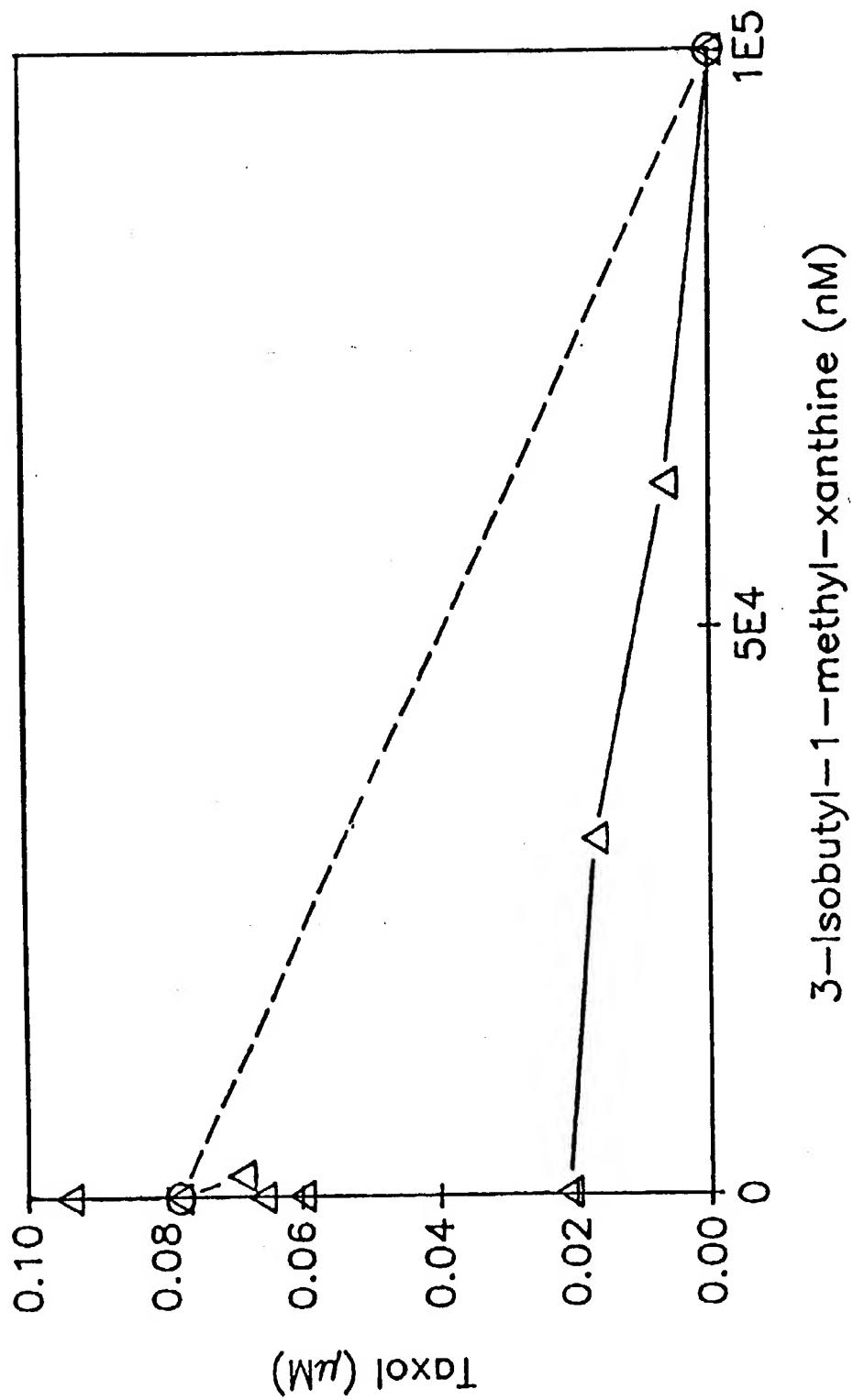
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FIGURE 11

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FIGURE 11.

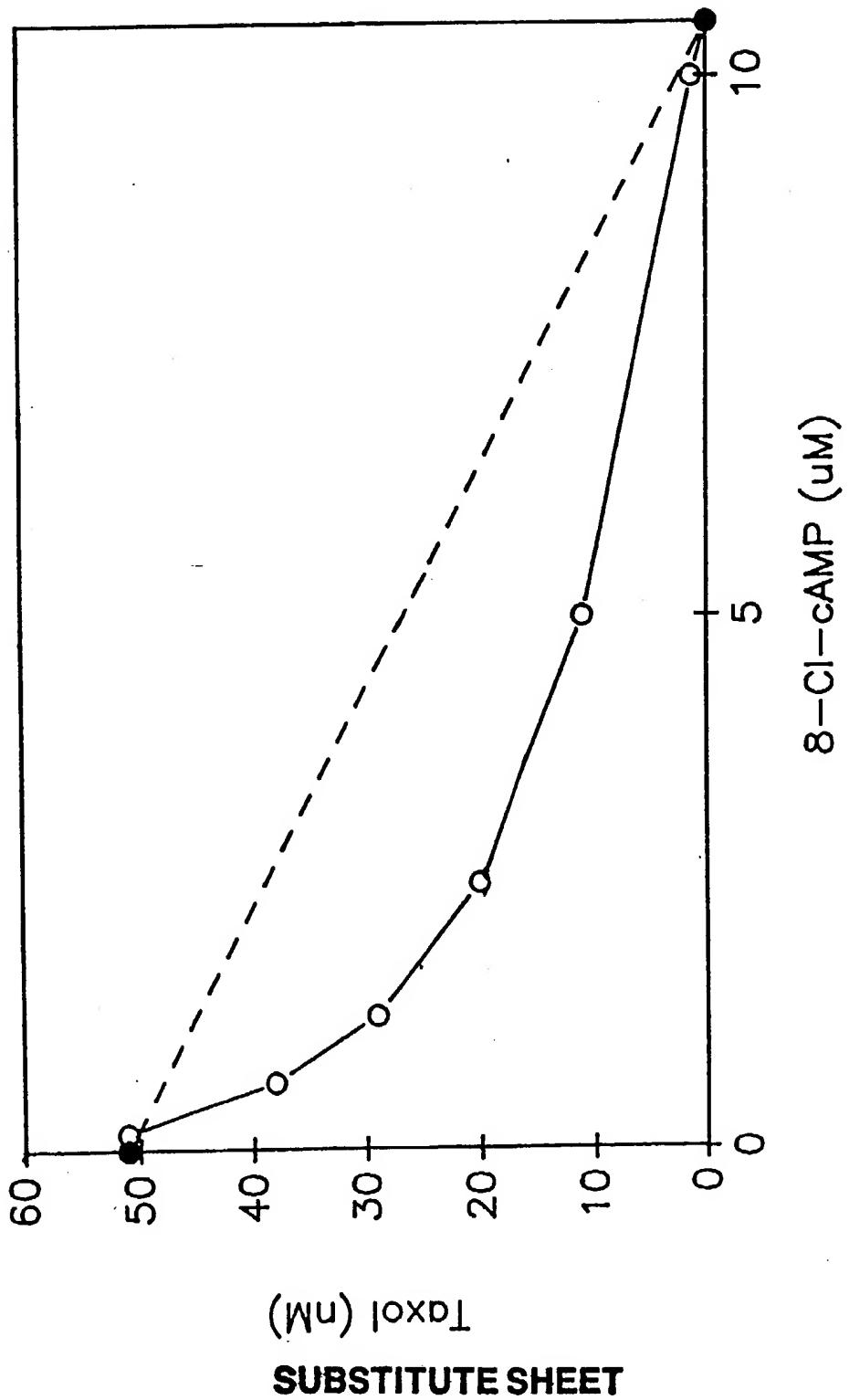
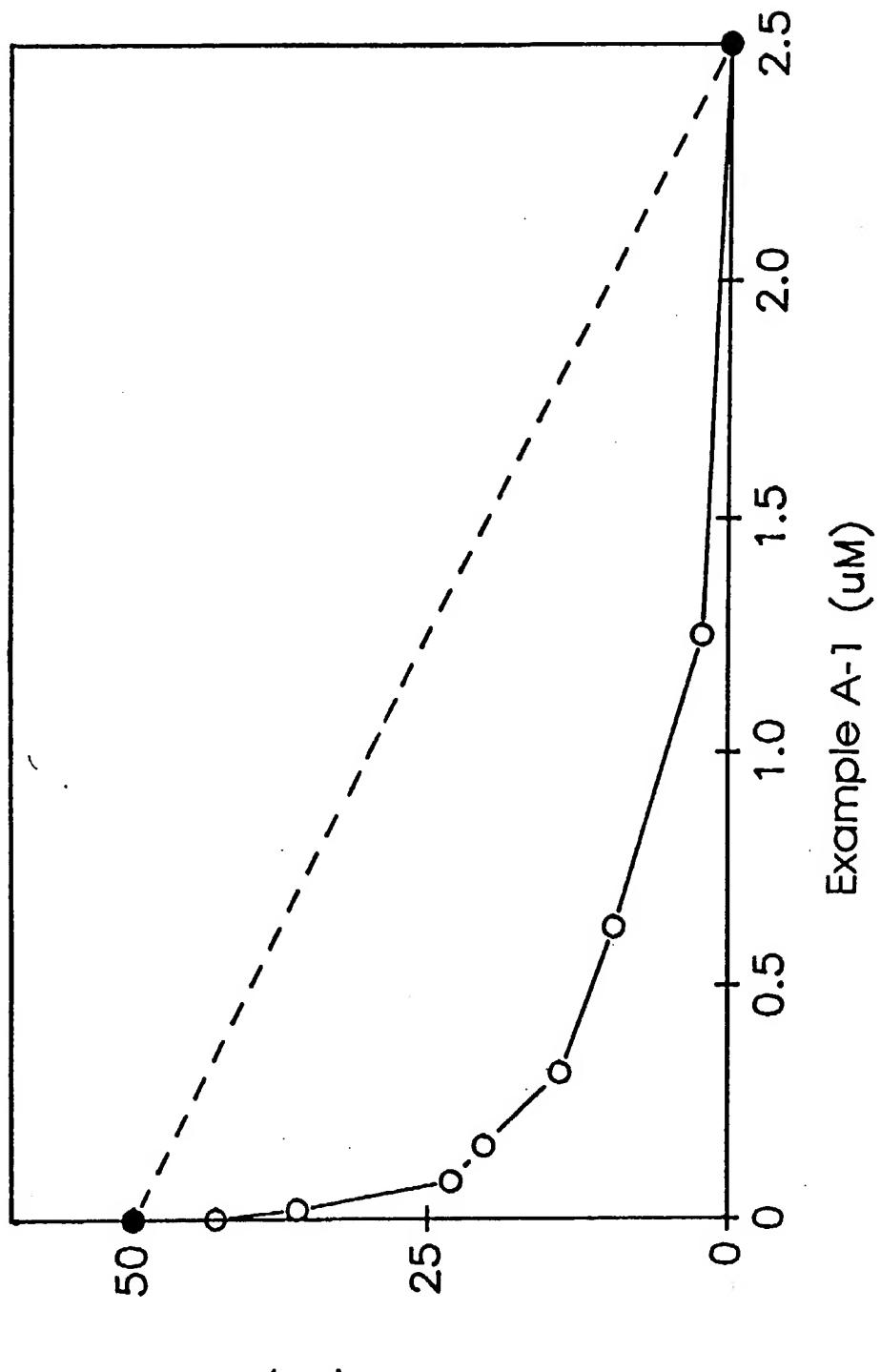
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FIGURE 12

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FIGURE 12.



Taxol (nM)

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FIGURE 13

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FIGURE 13.

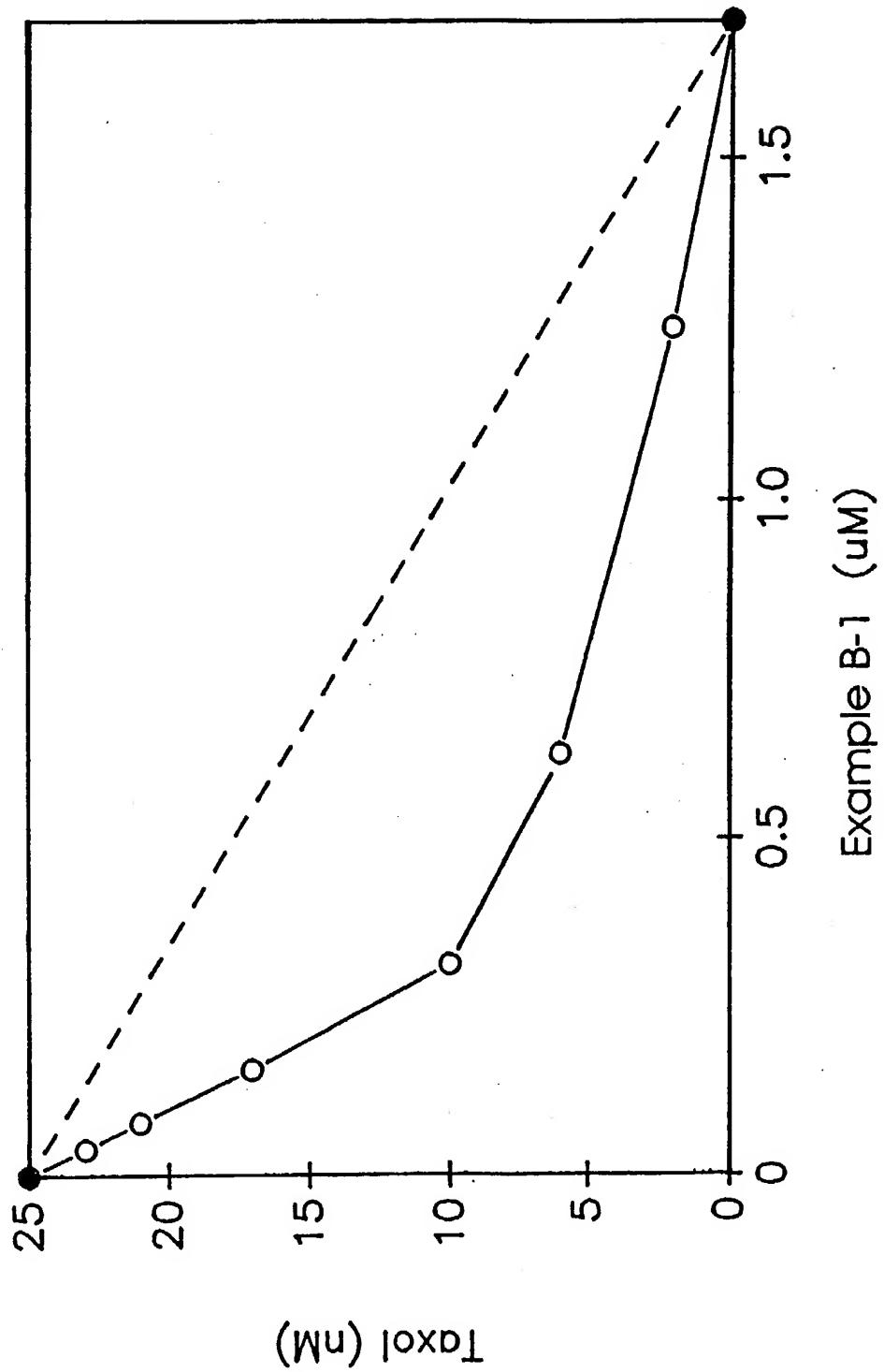
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FIGURE 14.

FIGURE 14

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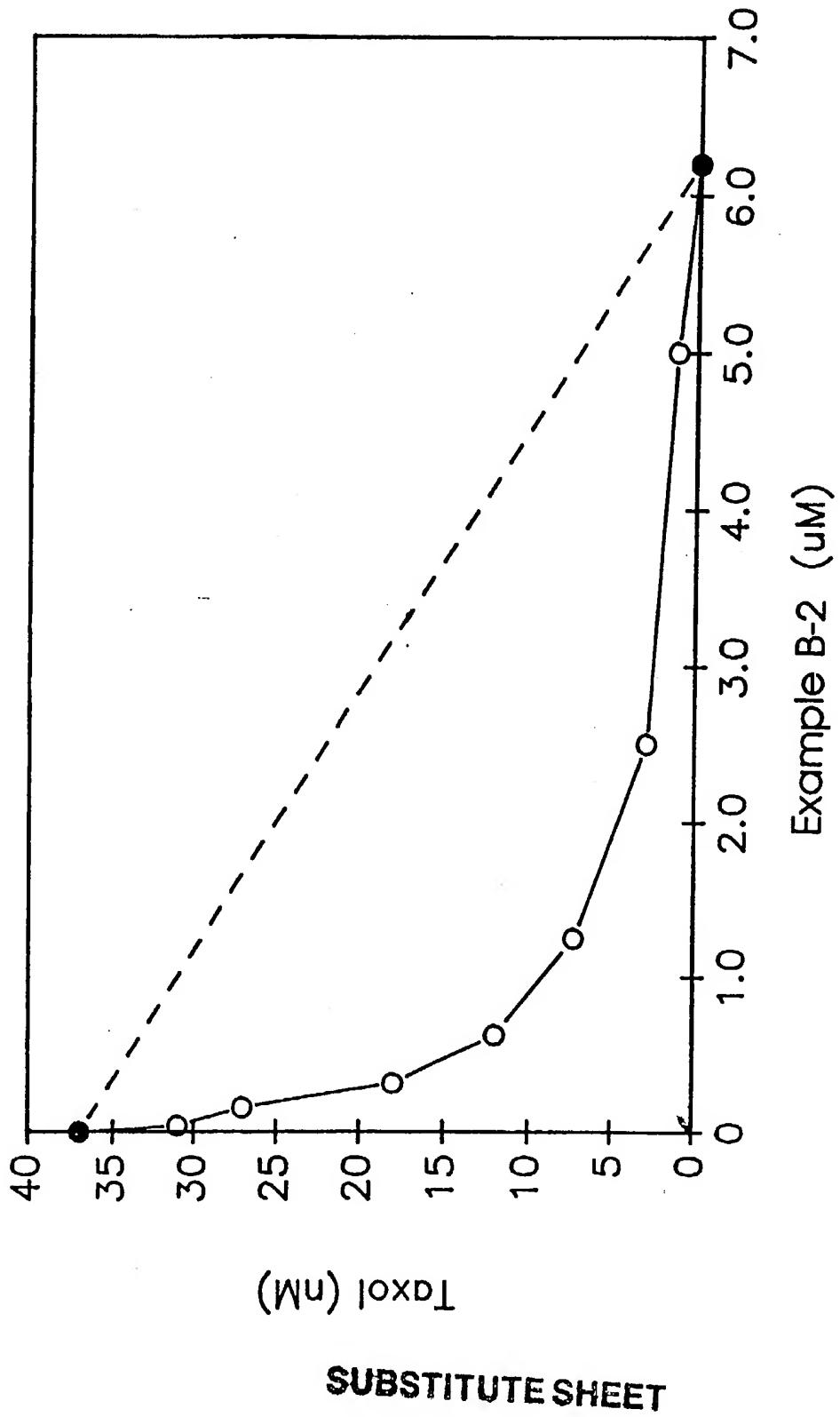


FIGURE 15

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FIGURE 15.

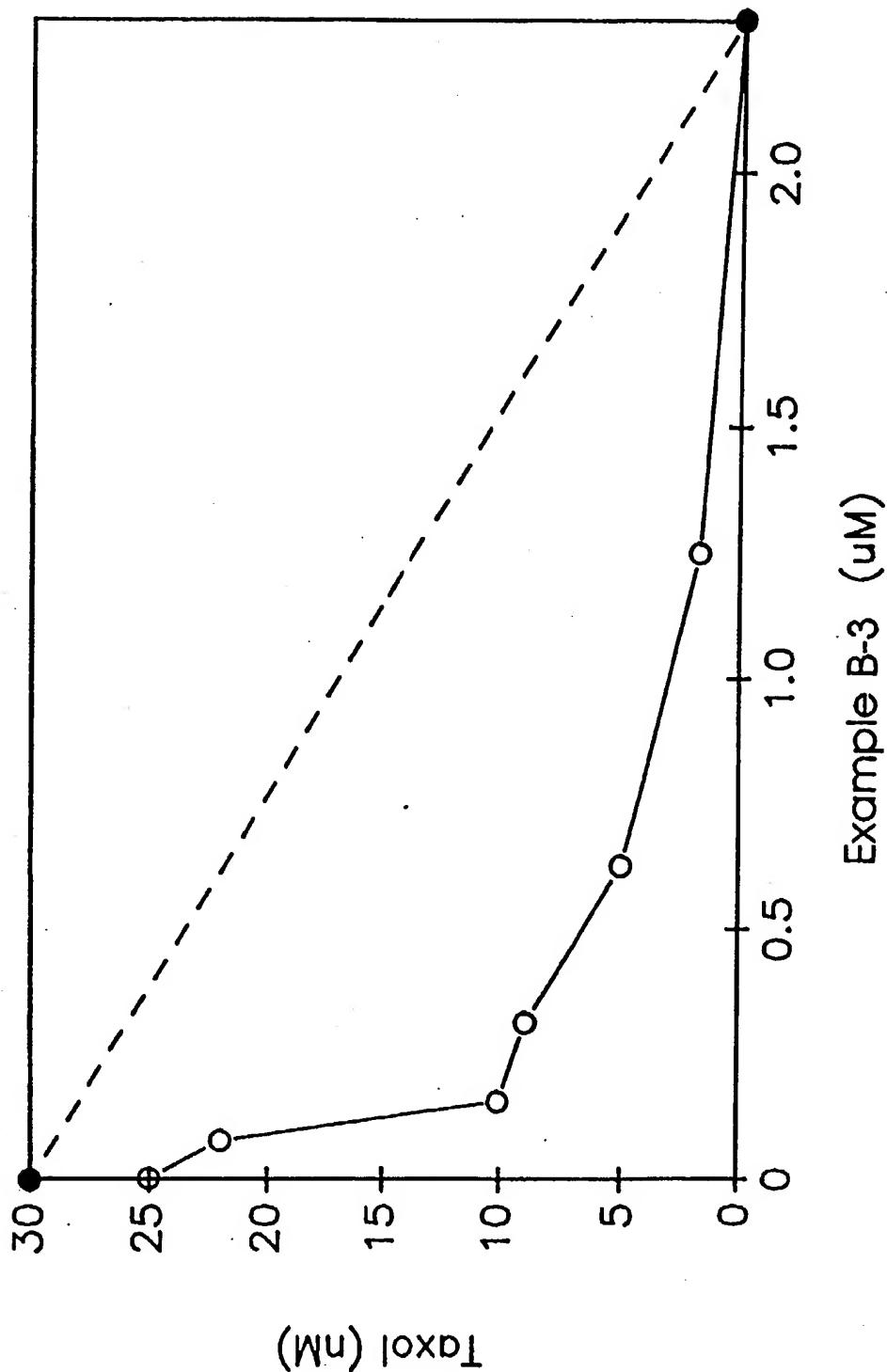
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FIGURE 16.

FIGURE 16

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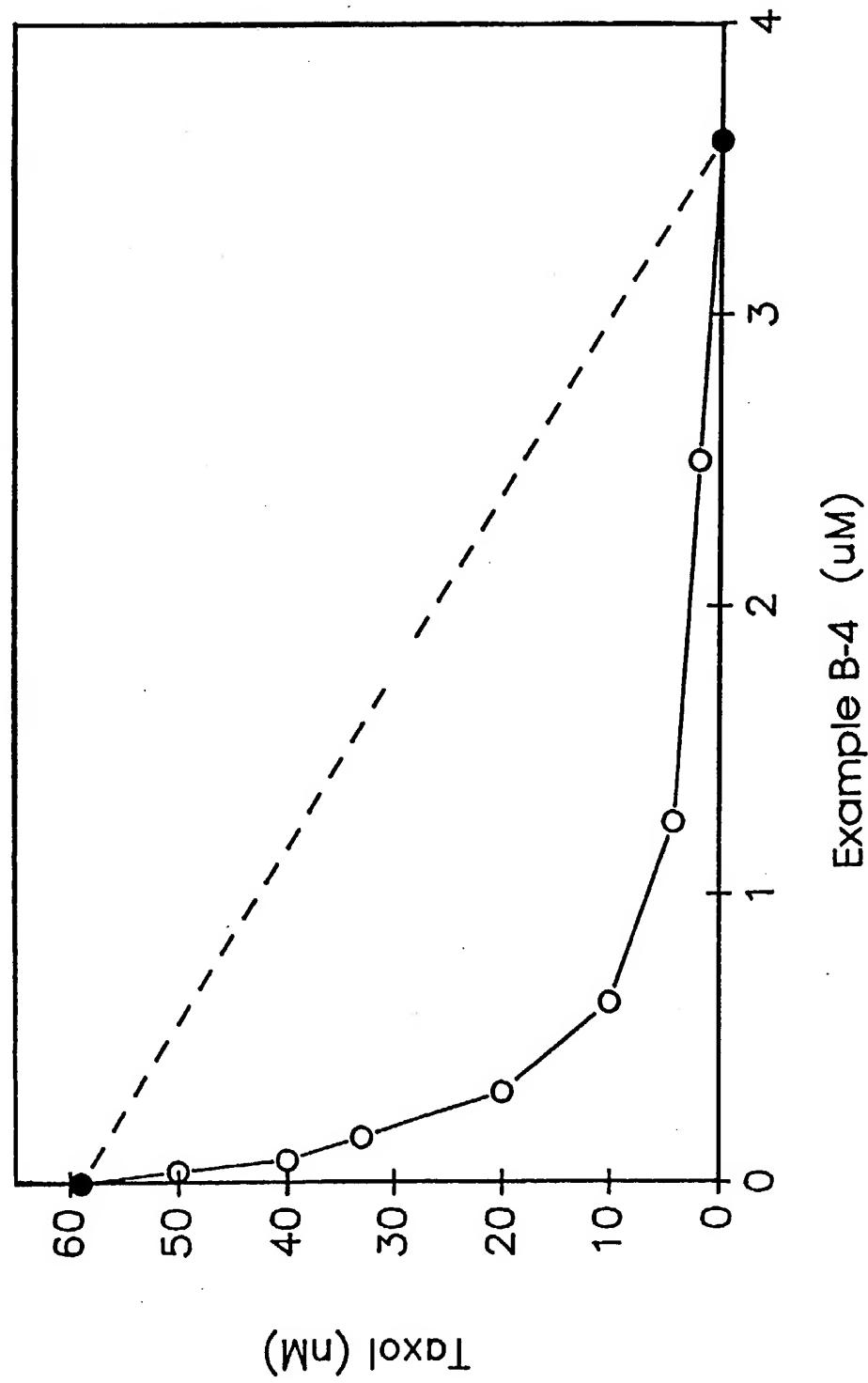
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FIGURE 17.

FIGURE 17

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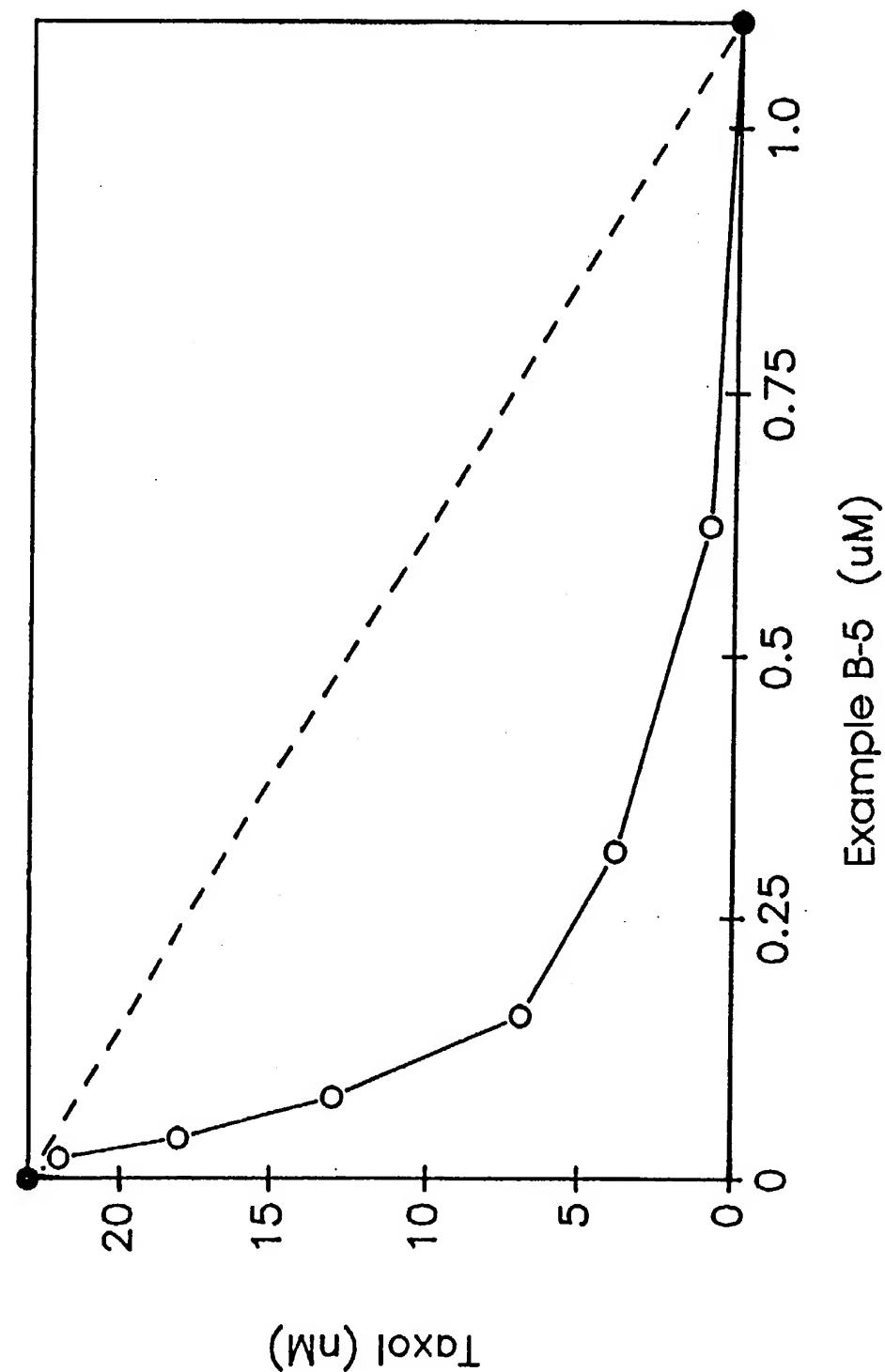
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FIGURE 18

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FIGURE 18.

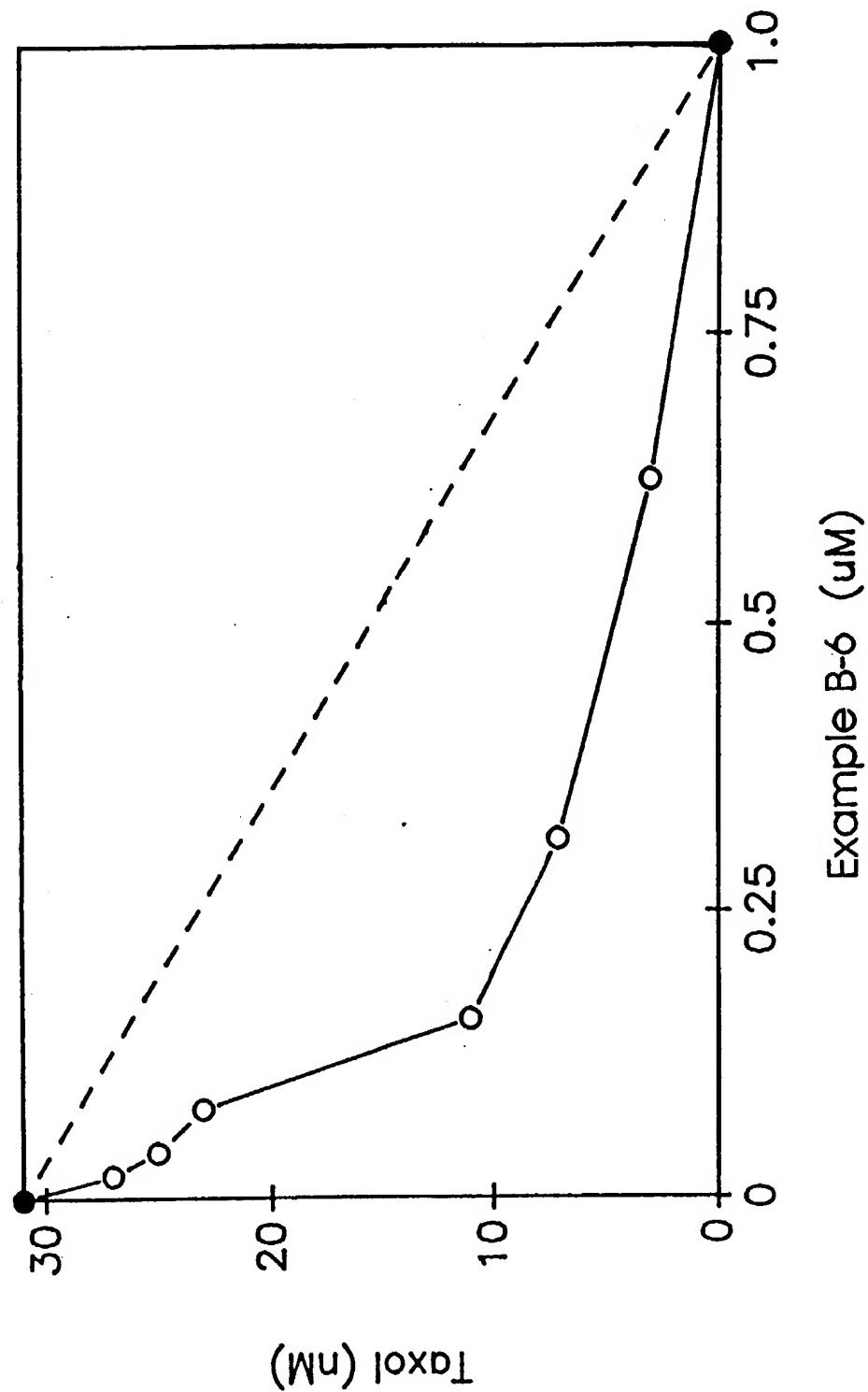
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FIGURE 19

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FIGURE 19.

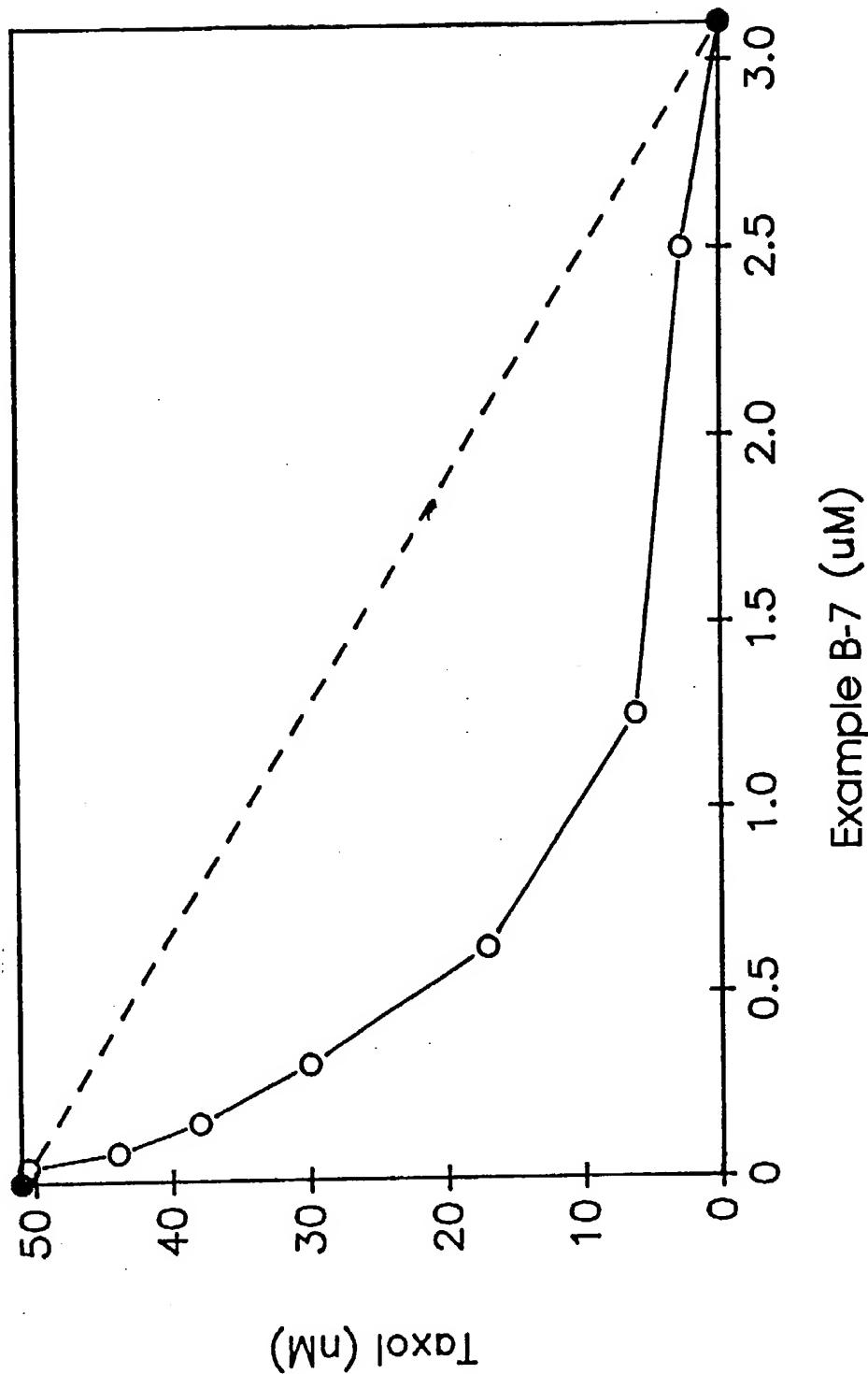
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FIGURE 20

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FIGURE 20.

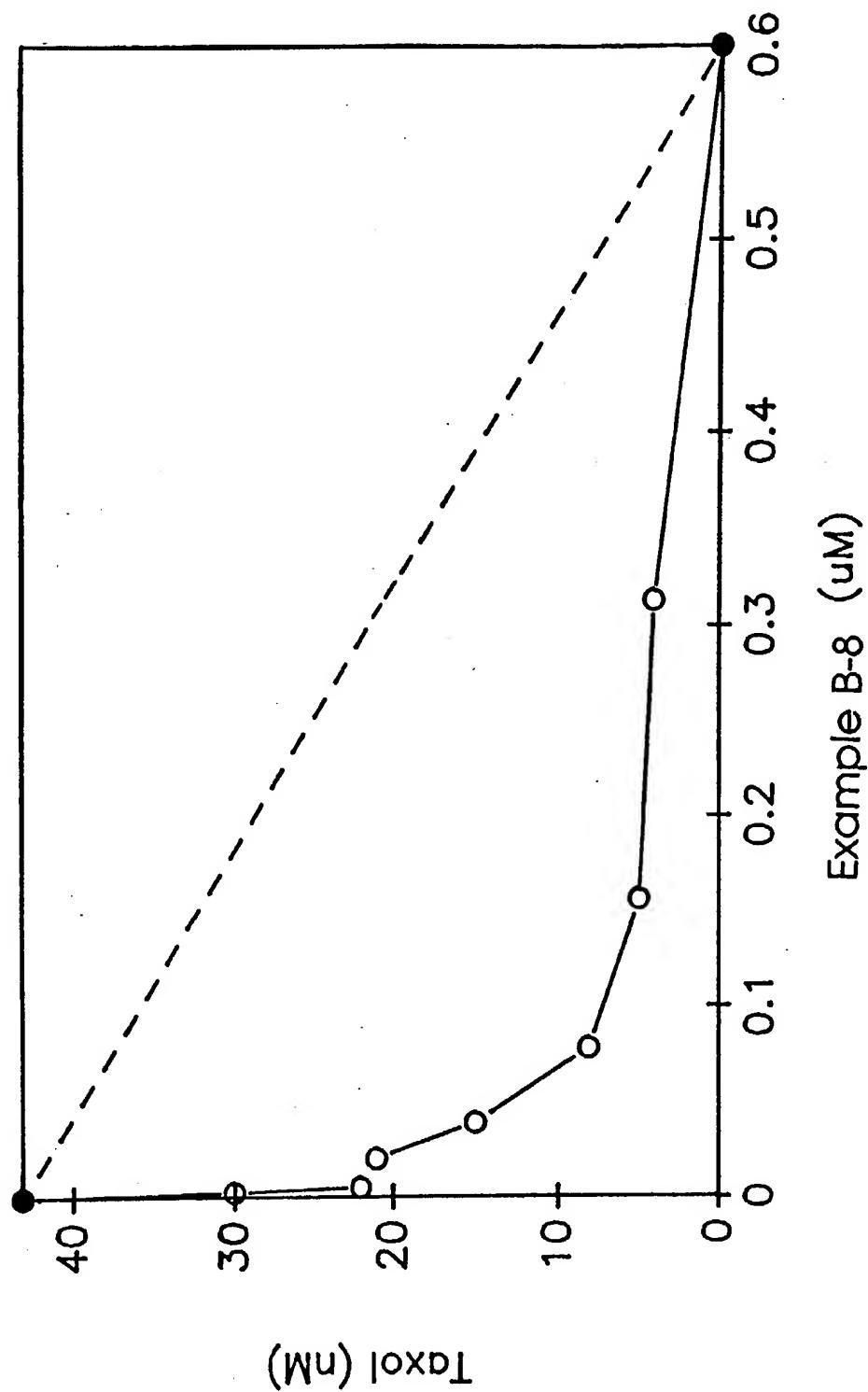
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FIGURE 21

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FIGURE 21.

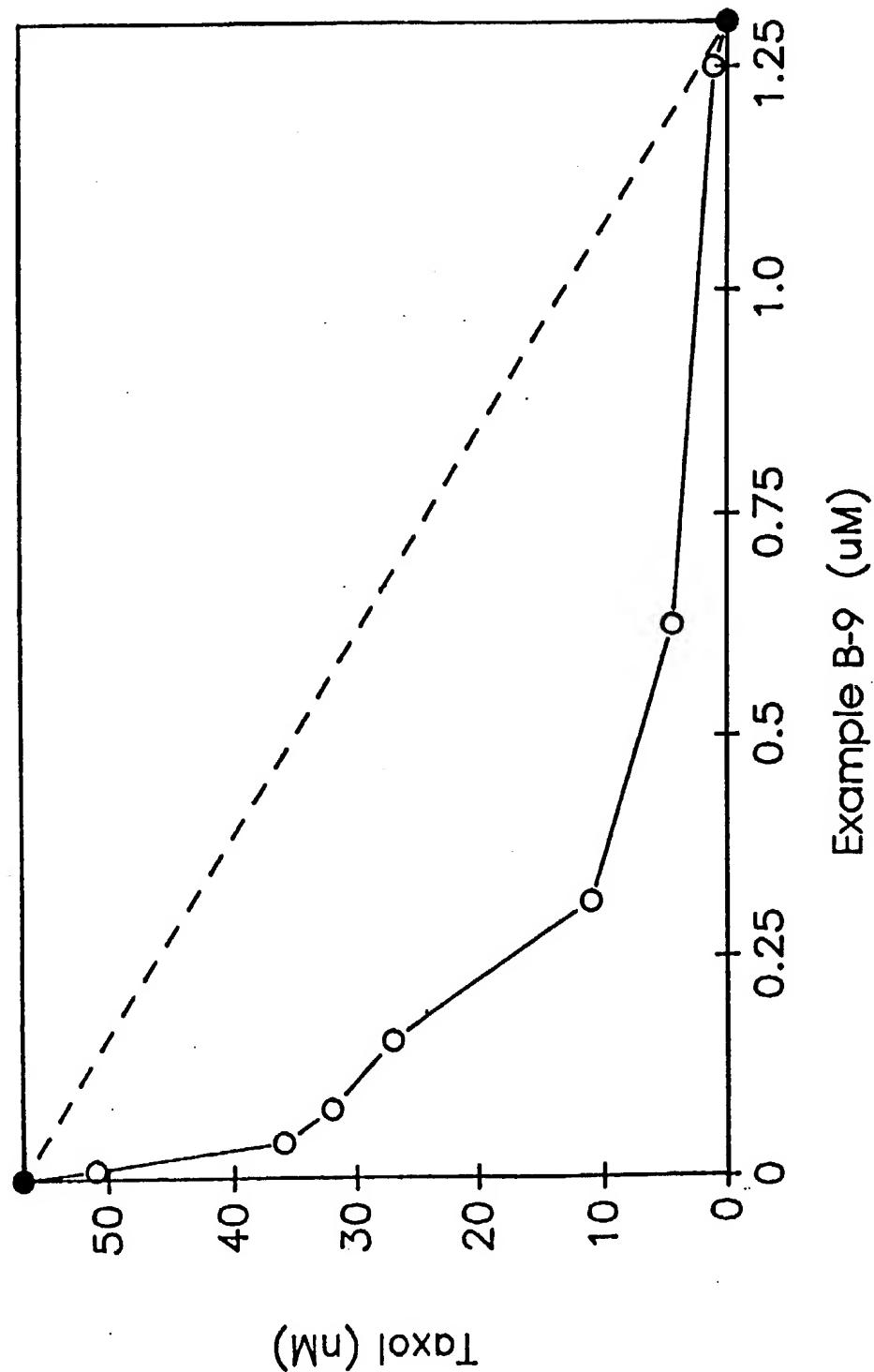
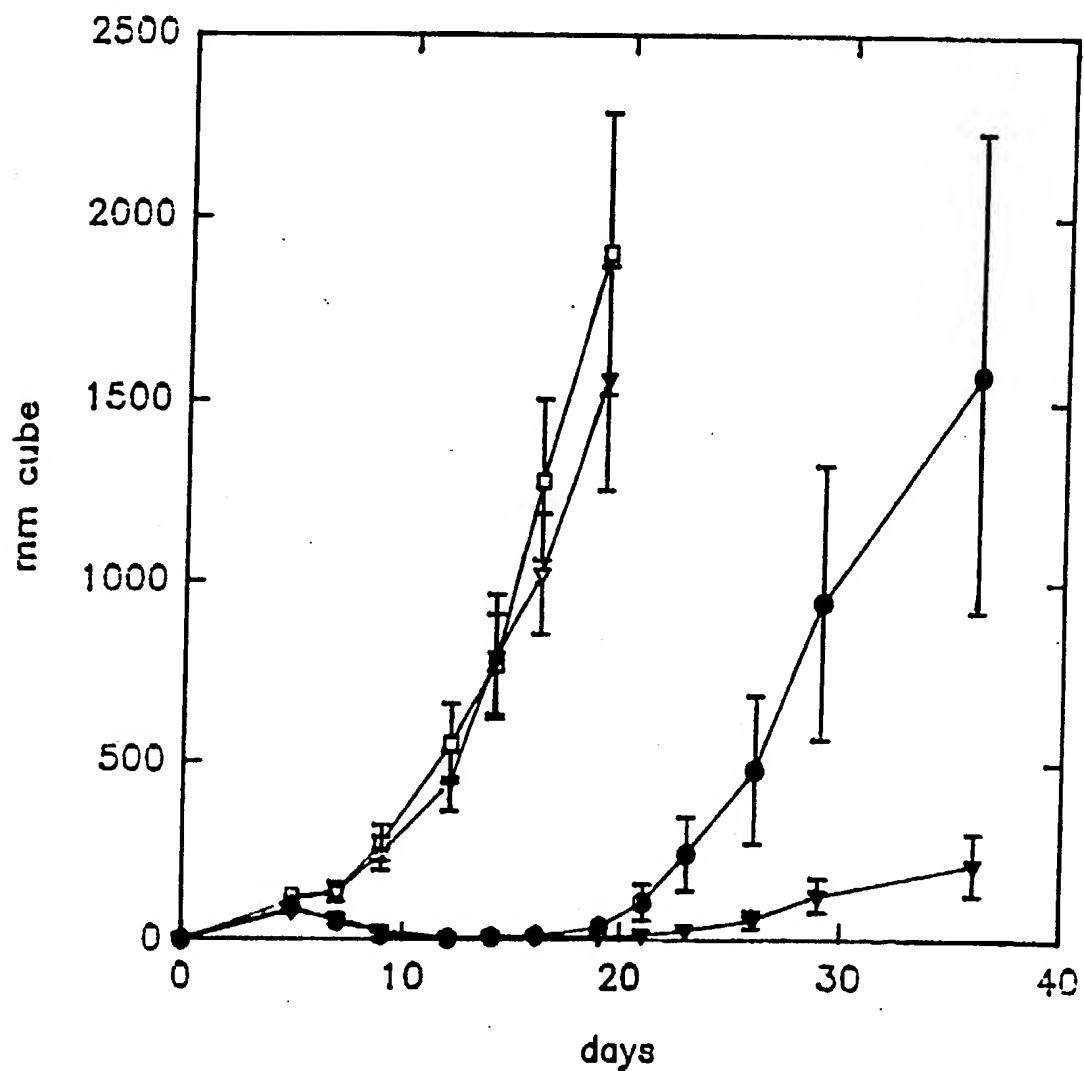
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FIGURE 22

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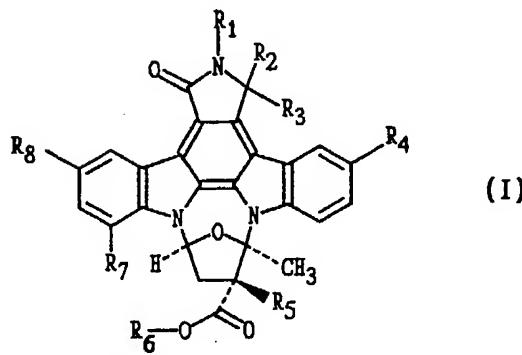
**FIGURE 22. TAXOL PLUS KT5720
ON TUMORED MICE****SUBSTITUTE SHEET**



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C07D 498/22, A61K 31/335, 31/55 // C07D 498/22, 307:00, 273:00 209:00, 209:00 209:00		A3	(11) International Publication Number: WO 94/04541
			(43) International Publication Date: 3 March 1994 (03.03.94)
(21) International Application Number: PCT/US93/07054			(72) Inventors; and
(22) International Filing Date: 30 July 1993 (30.07.93)			(75) Inventors/Applicants (for US only) : ABRAHAM, Irene [US/US]; 1805 Waite, Kalamazoo, MI 49008 (US), ARISTOFF, Paul, A. [US/US]; 1650 Brookmoor Lane, Portage, MI 49002 (US). SKULNICK, Harvey, I. [US/US]; 1745 Old Deer Run, Kalamazoo, MI 49009 (US).
(30) Priority data: 07/929,199 12 August 1992 (12.08.92) US			(74) Agent: WOOTTON, Thomas, A.; The Upjohn Company, 301 Henrietta Street, Kalamazoo, MI 49001 (US).
(60) Parent Application or Grant			(81) Designated States: AT, AU, BB, BG, BR, BY, CA, CH, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, KZ, LK, LU, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, US, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).
(63) Related by Continuation US Filed on 07/929,199 (CIP) 11 August 1992 (11.08.92)			Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.
(71) Applicant (for all designated States except US): THE UP-JOHN COMPANY [US/US]; 301 Henrietta Street, Kalamazoo, MI 49001 (US).			(88) Date of publication of the international search report: 09 June 1994 (09.06.94)

(54) Title: PROTEIN KINASE INHIBITORS AND RELATED COMPOUNDS COMBINED WITH TAXOL



(57) Abstract

This invention describes both known and novel compounds, some of which are protein kinase inhibitors, that may be combined with taxol type compounds. The combination of disclosed compounds plus taxol type compounds exhibits powerful synergistic effects and the combinations are useful in the treatment of cancer. The novel compounds and their synthesis are described. A compound of formula (I), above, is described wherein R₁-R₈ represent various substituents.

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 93/07054

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 5 C07D498/22 A61K31/335 A61K31/55 //C07D498/22, 307:00,
 273:00, 209:00, 209:00, 209:00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 5 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>CHEMICAL ABSTRACTS, vol. 111, no. 21, 20 November 1989, Columbus, Ohio, US; abstract no. 194456g, T. HIRATA ET AL 'Preparation of K-252 derivatives as protein kinase C inhibitors and formulations containing them' page 743 ; cited in the application see abstract & JP,A,63 295 589 (KYOWA HAKKO HOGYO CO) 1 December 1988</p> <p>---</p> <p>-/-</p>	1

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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- *'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *'&' document member of the same patent family

5 Date of the actual completion of the international search

11 January 1994

Date of mailing of the international search report

20.04.94

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INTERNATIONAL SEARCH REPORT

Intern: al Application No
PCT/US 93/07054

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	CHEMICAL ABSTRACTS, vol. 107, no. 25, 21 December 1987, Columbus, Ohio, US; abstract no. 236750y, T. HIRATA ET AL 'Preparation of k-252 derivatives as protein kinase C inhibitors and drugs' page 796 ; cited in the application see abstract & JP,A,62 155 284 (KYOWA HAKKO KOGYO CO) 10 July 1987 ---	1
A	EP,A,0 303 697 (KYOWA HAKKO KOGYO CO) 22 February 1989 cited in the application see claim 1 ---	1
A	EP,A,0 323 171 (KYOWA HAKKO KOGYO CO) 5 July 1989 cited in the application see claim 1 ---	1
A	CHEMICAL ABSTRACTS, vol. 103, no. 21, 25 November 1985, Columbus, Ohio, US; abstract no. 171551e, T. TSURUO ET AL 'Potentiation of antitumor agents by calcium channel blockers with special reference to cross-resistance patterns' page 29 ; see abstract & CANCER CHEMOTHER. PHARMACOL., vol.15, no.1, 1985 pages 16 - 19 ---	24,26
A	PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, vol.85, no.2, January 1988, WASHINGTON US pages 582 - 586 R. L. FINE ET AL 'Phorbol esters induce multidrug resistance in human breast cancer cells' see the whole document -----	24,26

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 93/07054

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
Claim 27 does not define the matter for which protection is sought (Art.6).
Claim not searched: 27

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

For further information please see Form PCT/ISA/206 mailed on 19.01.94

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-24, 26

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 93/07054

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
JP-A-63295589	01-12-88	NONE		
JP-A-62155284	10-07-87	JP-B-	5001794	11-01-93
EP-A-0303697	22-02-89	WO-A- US-A-	8807045 4923986	22-09-88 08-05-90
EP-A-0323171	05-07-89	JP-A- US-A-	1168689 4877776	04-07-89 31-10-89